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(54) Title: INHIBITORS OF HISTONE DEACETYLASE

(57) Abstract: Compounds having the formula or therapeutically acceptable salts thereof, are histone deacetylase (HDAC) inhibitors. Preparation of the compounds, compositions containing the compounds, and treatment of diseases using the compounds are disclosed.

INHIBITORS OF HISTONE DEACETYLASE

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Technical Field

10 The present invention relates to compounds which are useful for inhibiting histone deacetylase (HDAC), methods of making the compounds, compositions containing the compounds, and methods of treatment using the compounds.

Background of the Invention

15 The organized packing of DNA in the eukaryotic nucleus plays an important role in the regulation of gene transcription. DNA's highly condensed state is a consequence of its wrapping into chromatin. The fundamental repeating structural unit of chromatin is the nucleosome, which consists of 146 bases of DNA wrapped around a complex of eight histone proteins, two molecules each of the core histones, H2A, H2B, H3, and H4. Each core histone octomer is comprised of several highly conserved structural motifs including a globular 20 domain and an N-terminal tail domain that extends outside of the nucleosome. These histone N-terminal tails are enriched in basic amino acids, and are thought to mediate histone-DNA contacts through electrostatic interactions with DNA's negatively charged phosphate backbone. Based on the x-ray crystal structure of the nucleosome core particle, N-terminal histone tails also form contacts with the surface of histones of neighboring nucleosomes.

25 The capacity of histones to compact DNA is influenced by a number of post-translational modifications that occur on the N-terminal histone tails. One modification involves the reversible acetylation and deacetylation of the epsilon-amino group of lysine moieties found within the histone tails. The net level of acetylation of N-terminal histone tails is controlled by the activities of two families of enzymes, the histone acetyltransferases 30 (HATs) and histone deacetylases (HDACs). The identification of coactivator complexes that possess intrinsic HAT activity strongly supports the connection between histone acetylation and transcriptional activation (Bioessays 1998, 20, 615). Similarly, transcriptional repressor complexes have been shown to recruit HDACs to the promoter of target genes.

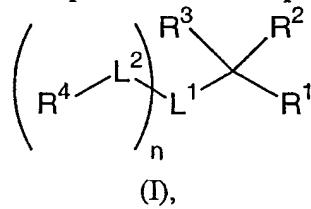
35 Several human cancers have been associated with malfunctions in HAT and HDAC activity. One example is the translocation of chromosomes 15 and 17 seen in the majority of acute promyelocytic leukemia patients. This translocation leads to the formation of a chimeric protein composed of the retinoic acid receptor fused to the PML transcription factor

(PML-RARA) (Mol. and Cell. Bio. 1998, 18, 7176). The recruitment of HDACs by this fusion protein diminishes its responsiveness to retinoic acid resulting in inhibition of differentiation of hematopoietic cells, one of the characteristic features of this disease.

Inhibition of the action of HDACs causes a variety of cellular responses including the
 5 accumulation of hyperacetylated histones, altered gene expression, and cell cycle arrest. Antiproliferative and antitumor properties have also been described for compounds possessing HDAC inhibitory activity (J. Biol. Chem. 1999, 274, 34940). While a number of natural product and synthetic HDAC inhibitors have been reported (J. Med. Chem. 1999, 42, 3001; and PNAS, 1998, 95, 3003), there still exists a need for inhibitors with improved
 10 profiles of activity.

Summary of the Invention

In its principle embodiment the present invention provides a compound of formula (I)



15

or a therapeutically acceptable salt thereof, wherein

n is 1 or 2;

20 L^1 is selected from the group consisting of alkenylene, alkylene, alkynylene, cycloalkylene, heteroalkylene, -(alkylene)-C(O)N(R^5)-(alkylene)-, -(alkylene)-O-(alkylene)-; wherein each group is drawn with its left-hand end being the end which attaches to L^2 , and its right-hand end being the end which attaches to the carbon substituted with R^1 , R^2 , and R^3 ;

25 L^2 is selected from the group consisting of a bond, C_2 alkenylene, -O-, -S-, -SO₂-, -OC(O)NR⁵-, -N(R^6)C(O)-, -C(O)N(R^6)-, -SO₂N(R^6)-, -N(R^6)SO₂-, -C=N-O-, -N(R^6)C(O)N(R^6)-, and -C(O)N(R^6)N(R^6)C(O)-; wherein each group is drawn with its left-hand end being the end which attaches to R^4 , and its right-hand end being the end which attaches to L^1 ;

30 R^1 is selected from the group consisting of alkanoyl, alkoxy carbonyl, aminocarbonyl, carboxy, haloalkyl, and heterocycle, wherein the heterocycle is selected from the group consisting of oxazolyl, dihydrooxazolyl, oxadiazolyl, and tetraazolyl;

R^2 and R^3 are hydroxy; or

R^2 and R^3 together are oxo;

35 R^4 is selected from the group consisting of alkoxyalkyl, alkyl, aryl, arylalkyl, cycloalkyl, (cycloalkyl)alkyl, heterocycle, and (heterocycle)alkyl; and

R^5 and R^6 are independently selected from the group consisting of hydrogen, alkyl,

35 aryl, and arylalkyl; or

R^4 and R^6 , together with the nitrogen atom to which they are attached, form a heterocycle selected from the group consisting of morpholinyl, piperazinyl, piperidinyl, and thiomorpholinyl; wherein the morpholinyl, the piperazinyl, the piperidinyl, and the thiomorpholinyl can be optionally substituted with one, two, or three substituents 5 independently selected from the group consisting of alkyl and spiroheterocycle.

In another embodiment, the present invention discloses a compound according to Claim 1 wherein n is 2.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R^1 is heterocycle, wherein the heterocycle is selected from the group 10 consisting of oxazolyl, dihydrooxazolyl, oxadiazolyl, and tetraazolyl; and L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R^1 is selected from the group consisting of alkoxy carbonyl and carboxy; and L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene.

15 In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R^1 is alkanoyl; and L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R^1 is aminocarbonyl; and L^1 is -(alkylene)-O-(alkylene)-.

20 In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R^1 is aminocarbonyl, L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene; and L^2 is selected from the group consisting of -O-, -S-, -SO₂-, and -SO₂N(R₆)-.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R^1 is aminocarbonyl, L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene; and L^2 is selected from the group consisting of -N(R⁶)C(O)N(R⁶)- and -C(O)N(R⁶)-.

25 In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R^1 is aminocarbonyl, L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene; and L^2 is selected from the group consisting of a bond, -C=N-O-, and -N(R⁶)C(O)CHC(O)N(R⁵)(R⁶)-.

30 In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R^1 is aminocarbonyl, L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene; and L^2 is -N(R⁶)C(O)-.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R^1 is aminocarbonyl, L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene; and L^2 is selected from the group consisting of -N(R⁶)C(O)N(R⁶)- and -C(O)N(R⁶)N(R⁶)C(O)-.

35 In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R^1 is haloalkyl; and L^1 is selected from the group consisting of alkenylene,

wherein the alkenylene is C₆ alkenylene; alkynylene, wherein the alkynylene is C₆ alkynylene; cycloalkylene; and -(alkylene)C(O)N(R⁵)(alkylene)-.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R¹ is haloalkyl; L¹ is alkylene, wherein the alkylene is C₅-C₇ alkylene; and L² is C₂ alkenylene.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R¹ is haloalkyl; L¹ is alkylene, wherein the alkylene is C₅-C₇ alkylene; and L² is -OC(O)N(R⁵)-.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R¹ is haloalkyl; L¹ is alkylene, wherein the alkylene is C₅-C₇ alkylene; and L² is -O-.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R¹ is haloalkyl; L¹ is alkylene, wherein the alkylene is C₅-C₇ alkylene; L² is -N(R⁶)C(O)-; and R⁴ is selected from the group consisting of alkoxyalkyl and alkyl.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R¹ is haloalkyl; L¹ is alkylene, wherein the alkylene is C₅-C₇ alkylene; L² is -N(R⁶)C(O)-; and R⁴ is arylalkyl.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R¹ is haloalkyl; L¹ is alkylene, wherein the alkylene is C₅-C₇ alkylene; L² is -N(R⁶)C(O)-; and R⁴ is arylalkyl.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R¹ is haloalkyl; L¹ is alkylene, wherein the alkylene is C₅-C₇ alkylene; L² is -N(R⁶)C(O)-; and R⁴ is selected from the group consisting of cycloalkyl, heterocycle, and (heterocycle)alkyl.

In another embodiment, the present invention provides a compound of formula (I) wherein n is 1; R¹ is haloalkyl; L¹ is alkylene, wherein the alkylene is C₅-C₇ alkylene; L² is -N(R⁶)C(O)-; and R⁴ and R⁶, together with the nitrogen atom to which they are attached, form a ring selected from the group consisting of morpholinyl, piperazinyl, piperidinyl, and thiomorpholinyl.

In another embodiment, the present invention provides a pharmaceutical composition comprising a compound of formula (I), or a therapeutically acceptable salt thereof, in combination with a pharmaceutically acceptable carrier.

In another embodiment, the present invention provides a method of inhibiting histone deacetylase in a patient in recognized need of such treatment comprising administering to the patient a therapeutically acceptable amount of a compound of formula (I), or a therapeutically acceptable salt thereof.

Detailed Description of the Invention

Compounds of the present invention are useful for the treatment of diseases in which histone deacetylase plays a role.

As used in the present specification the following terms have the meanings indicated:

5 The term "alkanoyl," as used herein, represents an alkyl group attached to the parent molecular moiety through a carbonyl group.

The term "alkenylene," as used herein, represents a divalent group of two to ten carbon atoms derived from a straight or branched chain hydrocarbon containing at least one double bond.

10 The term "C₂ alkenylene," as used herein, represents a divalent group of two carbon atoms containing a double bond.

The term "C₆ alkenylene," as used herein, represents a divalent group of six carbon atoms containing at least one double bond.

15 The term "alkoxy," as used herein, represents an alkyl group attached to the parent molecular moiety through an oxygen atom.

The term "alkoxyalkyl," as used herein, represents an alkoxy group attached to the parent molecular moiety through an alkyl group.

The term "alkoxycarbonyl," as used herein, represents an alkoxy group attached to the parent molecular moiety through a carbonyl group.

20 The term "alkyl," as used herein, represents a group of one to twelve carbon atoms derived from a straight or branched chain saturated hydrocarbon.

The term "alkylene," as used herein, represents a divalent group of one to ten carbon atoms derived from a straight or branched chain saturated hydrocarbon. The alkylene groups of the present invention can be optionally substituted with a hydroxy group.

25 The term "C₅-C₇ alkylene," as used herein, represents a divalent group of five to seven carbon atoms derived from a straight or branched chain saturated hydrocarbon. The C₅-C₇ alkylene groups of the present invention can be optionally substituted with a hydroxy group.

30 The term "C₆ alkylene," as used herein, represents a divalent group of six carbon atoms derived from a straight or branched chain saturated hydrocarbon. The C₆ alkylene groups of the present invention can be optionally substituted with a hydroxy group.

The term "alkylsulfanyl," as used herein, represents an alkyl group attached to the parent molecular moiety through a sulfur atom.

35 The term "alkylsulfonyl," as used herein, represents an alkyl group attached to the parent molecular moiety through a sulfonyl group.

The term "alkynylene," as used herein, represents a divalent group of two to ten carbon atoms derived from a straight or branched chain hydrocarbon containing at least one

triple bond.

The term "C₆ alkynylene," as used herein, represents a divalent group of six carbon atoms derived from a straight or branched chain hydrocarbon containing at least one triple bond.

5 The term "amino," as used herein, represents -NR⁷R⁸, wherein R⁷ and R⁸ are independently selected from the group consisting of hydrogen, alkanoyl, alkyl, cycloalkyl, (cycloalkyl)alkyl, a nitrogen protecting group, and unsubstituted aryl.

The term "aminocarbonyl," as used herein, represents an amino group attached to the parent molecular moiety through a carbonyl group.

10 The term "aryl," as used herein, represents a phenyl group or a bicyclic or tricyclic fused ring system wherein one or more of the fused rings is a phenyl group. Bicyclic fused ring systems are exemplified by a phenyl group fused to a cycloalkyl group as defined herein, a cycloalkenyl group as defined herein, or another phenyl group. Tricyclic fused ring systems are exemplified by a bicyclic fused ring system fused to a cycloalkyl group as

15 defined herein, a cycloalkenyl group as defined herein, or another phenyl group.

Representative examples of aryl include, but are not limited to, anthracenyl, azulenyl, fluorenyl, indanyl, indenyl, naphthyl, phenyl, and tetrahydronaphthyl. Aryl groups having an unsaturated or partially saturated ring fused to an aromatic ring can be attached through the saturated or the unsaturated part of the group. The aryl groups of the present invention can be 20 optionally substituted with one, two, three, four, or five substituents independently selected from the group consisting of alkanoyl, alkoxy, alkyl, alkylsulfanyl, alkylsulfonyl, amino, aminoalkoxy, a second aryl, arylalkoxy, arylalkyl, arylcarbonyl, aryloxy, arylsulfanyl, arylsulfonyl, carbonyloxy, cyano, cycloalkyl, (cycloalkyl)alkyl, halo, haloalkoxy, haloalkyl, heterocycle, (heterocycle)alkoxy, (heterocycle)alkyl, hydroxy, nitro, and oxo; wherein the 25 second aryl, the aryl part of the arylalkoxy, the arylalkyl, the arylcarbonyl, the aryloxy, the arylsulfanyl, and the arylsulfonyl; the heterocycle; and the heterocycle part of the (heterocycle)alkyl can be further optionally substituted with one, two, three, four, or five substituents independently selected from the group consisting of alkoxy, alkyl, alkylsulfanyl, alkylsulfonyl, amino, cyano, halo, haloalkoxy, haloalkyl, hydroxy, and nitro.

30 The term "arylalkoxy," as used herein, represents an aryl group attached to the parent molecular moiety through an alkoxy group.

The term "arylalkyl," as used herein, represents an aryl group attached to the parent molecular moiety through an alkyl group. The alkyl part of the arylalkyl groups of the present invention can be optionally substituted with one or two substituents independently 35 selected from the group consisting of aminocarbonyl and aryl.

The term "arylcarbonyl," as used herein, represents an aryl group attached to the parent molecular moiety through a carbonyl group.

The term "aryloxy," as used herein, represents an aryl group attached to the parent molecular group through an oxygen atom.

The term "arylsulfanyl," as used herein, represents an aryl group attached to the parent molecular moiety through a sulfur atom.

5 The term "arylsulfonyl," as used herein, represents an aryl group attached to the parent molecular moiety through a sulfonyl group.

The term "carbonyl," as used herein, represents -C(O)-.

The term "carbonyloxy," as used herein, represents an alkanoyl group attached to the parent molecular moiety through an oxygen atom.

10 The term "carboxy," as used herein, represents -CO₂H.

The term "cyano," as used herein, represents -CN.

15 The term "cycloalkenyl," as used herein, represents a non-aromatic ring system having three to ten carbon atoms and one to three rings, wherein each five-membered ring has one double bond, each six-membered ring has one or two double bonds, each seven- and eight-membered ring has one to three double bonds, and each nine-to ten-membered ring has one to four double bonds. Examples of cycloalkenyl groups include cyclohexenyl, octahydronaphthalenyl, norbornenyl, and the like.

20 The term "cycloalkyl," as used herein, represents a saturated monocyclic, bicyclic, or tricyclic hydrocarbon ring system having three to twelve carbon atoms. Examples of cycloalkyl groups include cyclopropyl, cyclopentyl, bicyclo(3.1.1)heptyl, adamantyl, and the like.

The term "(cycloalkyl)alkyl," as used herein, represents a cycloalkyl group attached to the parent molecular moiety through an alkyl group.

25 The term "cycloalkylene," as used herein represents a divalent group derived from a saturated monocyclic hydrocarbon ring system having three to twelve carbon atoms.

The term "halo," or "halogen," as used herein, represents F, Cl, Br, or I.

The term "haloalkoxy," as used herein, represents a haloalkyl group attached to the parent molecular group through an oxygen atom.

30 The term "haloalkyl," as used herein, represents an alkyl group substituted by one, two, three, or four halogen atoms.

The term "heteroalkylene," as used herein, represents a divalent group of two to eight atoms derived from a saturated straight or branched chain containing one or two heteroatoms independently selected from the group consisting of nitrogen, oxygen, and sulfur, wherein the remaining atoms are carbon. The heteroalkylene groups of the present invention can be attached through the carbon atoms or the heteroatoms in the chain.

35 The term "heterocycle," as used herein, represents a monocyclic, bicyclic, or tricyclic ring system wherein one or more rings is a four-, five-, six-, or seven-membered ring

containing one, two, or three heteroatoms independently selected from the group consisting of nitrogen, oxygen, and sulfur. Monocyclic ring systems are exemplified by any 3- or 4-membered ring containing a heteroatom independently selected from the group consisting of oxygen, nitrogen and sulfur; or a 5-, 6- or 7-membered ring containing one, two or three heteroatoms wherein the heteroatoms are independently selected from the group consisting of nitrogen, oxygen and sulfur. The 3- and 4-membered rings have no double bonds, the 5-membered ring has from 0-2 double bonds and the 6- and 7-membered rings have from 0-3 double bonds. Representative examples of monocyclic ring systems include, but are not limited to, azetidine, azepine, aziridine, diazepine, 1,3-dioxolane, dioxane, dithiane, furan, 10 imidazole, imidazoline, imidazolidine, isothiazole, isothiazoline, isothiazolidine, isoxazole, isoxazoline, isoxazolidine, morpholine, oxadiazole, oxadiazoline, oxadiazolidine, oxazole, oxazoline, oxazolidine, piperazine, piperidine, pyran, pyrazine, pyrazole, pyrazoline, pyrazolidine, pyridine, pyrimidine, pyridazine, pyrrole, pyrroline, pyrrolidine, tetrahydrofuran, tetrahydrothiophene, tetrazine, tetrazole, thiadiazole, thiadiazoline, 15 thiadiazolidine, thiazole, thiazoline, thiazolidine, thiophene, thiomorpholine, thiomorpholine sulfone, thiopyran, triazine, triazole, trithiane, and the like. Bicyclic ring systems are exemplified by any of the above monocyclic ring systems fused to an aryl group as defined herein, a cycloalkyl group as defined herein, a cycloalkenyl group, as defined herein, or another monocyclic heterocycle ring system. Representative examples of bicyclic ring 20 systems include but are not limited to, benzimidazole, benzothiazole, benzothiophene, benzoxazole, benzofuran, benzopyran, benzothiopyran, benzodioxine, 1,3-benzodioxole, cinnoline, indazole, indole, indoline, indolizine, naphthyridine, isobenzofuran, isobenzothiophene, isoindole, isoindoline, isoquinoline, phthalazine, pyranopyridine, quinoline, quinolizine, quinoxaline, quinazoline, tetrahydroisoquinoline, tetrahydroquinoline, 25 thiopyranopyridine, and the like. Tricyclic rings systems are exemplified by any of the above bicyclic ring systems fused to an aryl group as defined herein, a cycloalkyl group as defined herein, a cycloalkenyl group as defined herein, or another monocyclic heterocycle ring system. Representative examples of tricyclic ring systems include, but are not limited to, acridine, carbazole, carboline, dibenzofuran, dibenzothiophene, naphthofuran, 30 naphthothiophene, oxanthrene, phenazine, phenoxathiin, phenoxyazine, phenothiazine, thianthrene, thioxanthene, xanthene, and the like. Heterocycle groups can be attached to the parent molecular moiety through a carbon atom or a nitrogen atom in the ring.

The heterocycle groups of the present invention can be optionally substituted with one, two, three, or four substituents independently selected from the group consisting of 35 alkanoyl, alkoxy, alkyl, alkylsulfanyl, amino, aryl, arylalkoxy, arylalkyl, arylcarbonyl, aryloxy, arylsulfanyl, carbonyloxy, cyano, cycloalkyl, (cycloalkyl)alkyl, halo, haloalkoxy, haloalkyl, a second heterocycle, hydroxy, nitro, oxo, and spiroheterocycle, wherein the

second heterocycle can be further optionally substituted with one, two, three, four, or five substituents independently selected from the group consisting of alkoxy, alkyl, amino, aminoalkoxy, cyano, halo, haloalkoxy, haloalkyl, a third heterocycle, hydroxy, and nitro. The third heterocycle can be further optionally substituted with one, two, or three substituents 5 independently selected from the group consisting of alkoxy, alkyl, cyano, halo, haloalkoxy, haloalkyl, hydroxy, nitro, and oxo.

The term "(heterocycle)alkoxy," as used herein, represents a heterocycle group attached to the parent molecular moiety through an alkoxy group.

10 The term "(heterocycle)alkyl," as used herein, represents a heterocycle group attached to the parent molecular group through an alkyl group.

The term "hydroxy," as used herein, represents -OH.

The term "nitro," as used herein, represents -NO₂.

15 The term "nitrogen protecting group," as used herein, represents groups intended to protect an amino group against undesirable reactions during synthetic procedures. Common N-protecting groups comprise acyl groups such as acetyl, benzoyl, 2-bromoacetyl, 4-bromobenzoyl, tert-butyloxycarbonyl, carboxaldehyde, 2-chloroacetyl, 4-chlorobenzoyl, α -chlorobutyryl, 4-nitrobenzoyl, o-nitrophenoxyacetyl, phthalyl, pivaloyl, propionyl, trichloroacetyl, and trifluoroacetyl; sulfonyl groups such as benzenesulfonyl, and p-toluenesulfonyl; carbamate forming groups such as benzyloxycarbonyl, benzyloxycarbonyl 20 (Cbz), tert-butyloxycarbonyl (Boc), p-chlorobenzyloxycarbonyl, p-methoxybenzyloxycarbonyl, and the like.

The term "oxo," as used herein, represents (=O).

25 The term "spiroheterocycle," as used herein, represents a heteroalkylene diradical, each end of which is attached to the same carbon atom of the parent molecular moiety. Examples of spiroheterocycles include dioxolanyl, tetrahydrofuranyl, pyrrolidinyl, and the like.

The term "sulfonyl," as used herein, represents -SO₂⁻.

30 The present compounds can also exist as therapeutically acceptable prodrugs. The term "therapeutically acceptable prodrug," refers to those prodrugs or zwitterions which are suitable for use in contact with the tissues of patients without undue toxicity, irritation, and allergic response, are commensurate with a reasonable benefit/risk ratio, and are effective for their intended use. The term "prodrug," refers to compounds which are rapidly transformed in vivo to parent compounds of formula (I) for example, by hydrolysis in blood.

35 The compounds of the present invention can exist as therapeutically acceptable salts. The term "therapeutically acceptable salt," as used herein, represents salts or zwitterionic forms of the compounds of the present invention which are water or oil-soluble or dispersible, which are suitable for treatment of diseases without undue toxicity, irritation, and

allergic response; which are commensurate with a reasonable benefit/risk ratio, and which are effective for their intended use. The salts can be prepared during the final isolation and purification of the compounds or separately by reacting an amino group with a suitable acid. Representative acid addition salts include acetate, adipate, alginic acid, citrate, aspartate, 5 benzoate, benzenesulfonate, bisulfate, butyrate, camphorate, camphorsulfonate, digluconate, glycerophosphate, hemisulfate, heptanoate, hexanoate, formate, fumarate, hydrochloride, hydrobromide, hydroiodide, 2-hydroxyethansulfonate, lactate, maleate, mesitylenesulfonate, methanesulfonate, naphthalenesulfonate, nicotinate, 2-naphthalenesulfonate, oxalate, pamoate, pectinate, persulfate, 3-phenylpropionate, picrate, pivalate, propionate, succinate, 10 tartrate, trichloroacetate, trifluoroacetate, phosphate, glutamate, bicarbonate, para-toluenesulfonate, and undecanoate. Also, amino groups in the compounds of the present invention can be quaternized with methyl, ethyl, propyl, and butyl chlorides, bromides, and iodides; dimethyl, diethyl, dibutyl, and diethyl sulfates; decyl, lauryl, myristyl, and steryl chlorides, bromides, and iodides; and benzyl and phenethyl bromides. Examples of acids 15 which can be employed to form therapeutically acceptable addition salts include inorganic acids such as hydrochloric, hydrobromic, sulfuric, and phosphoric, and organic acids such as oxalic, maleic, succinic, and citric.

Basic addition salts can be prepared during the final isolation and purification of the compounds by reacting a carboxy group with a suitable base such as the hydroxide, 20 carbonate, or bicarbonate of a metal cation or with ammonia or an organic primary, secondary, or tertiary amine. The cations of therapeutically acceptable salts include lithium, sodium, potassium, calcium, magnesium, and aluminum, as well as nontoxic quaternary amine cations such as ammonium, tetramethylammonium, tetraethylammonium, methylamine, dimethylamine, trimethylamine, triethylamine, diethylamine, ethylamine, 25 tributylamine, pyridine, N,N-dimethylaniline, N-methylpiperidine, N-methylmorpholine, dicyclohexylamine, procaine, dibenzylamine, N,N-dibenzylphenethylamine, 1-ephedamine, and N,N'-dibenzylethylenediamine. Other representative organic amines useful for the formation of base addition salts include ethylenediamine, ethanolamine, diethanolamine, piperidine, and piperazine.

30 In addition to the compounds of the present invention and their pharmaceutically acceptable salts, the invention is further directed, where applicable, to unsolvated as well as solvated forms of the compounds (e.g., hydrated forms) having the ability to inhibit HDAC.

Because carbon-carbon double bonds exist in the present compounds, the invention contemplates various geometric isomers and mixtures thereof resulting from the arrangement 35 of substituents around these carbon-carbon double bonds. It should be understood that the invention encompasses both isomeric forms, or mixtures thereof, which possess the ability to inhibit histone deacetylase. These substituents are designated as being in the E or Z

configuration wherein the term "E" represents higher order substituents on opposite sides of the carbon-carbon double bond, and the term "Z" represents higher order substituents on the same side of the carbon-carbon double bond.

In accordance with methods of treatment and pharmaceutical compositions of the invention, the compounds can be administered alone or in combination with other HDAC inhibitors. When using the compounds, the specific therapeutically effective dose level for any particular patient will depend upon factors such as the disorder being treated and the severity of the disorder; the activity of the particular compound used; the specific composition employed; the age, body weight, general health, sex, and diet of the patient; the time of administration; the route of administration; the rate of excretion of the compound employed; the duration of treatment; and drugs used in combination with or coincidentally with the compound used. The compounds can be administered orally, parenterally, osmotically (nasal sprays), rectally, vaginally, or topically in unit dosage formulations containing carriers, adjuvants, diluents, vehicles, or combinations thereof. The term "parenteral" includes infusion as well as subcutaneous, intravenous, intramuscular, and intrasternal injection.

Parenterally administered aqueous or oleaginous suspensions of the compounds can be formulated with dispersing, wetting, or suspending agents. The injectable preparation can also be an injectable solution or suspension in a diluent or solvent. Among the acceptable diluents or solvents employed are water, saline, Ringer's solution, buffers, monoglycerides, diglycerides, fatty acids such as oleic acid, and fixed oils such as monoglycerides or diglycerides.

The inhibitory effect of parenterally administered compounds can be prolonged by slowing their absorption. One way to slow the absorption of a particular compound is by administering injectable depot forms comprising suspensions of crystalline, amorphous, or otherwise water-insoluble forms of the compound. The rate of absorption of the compound is dependent on its rate of dissolution which is, in turn, dependent on its physical state. Another way to slow absorption of a particular compound is by administering injectable depot forms comprising the compound as an oleaginous solution or suspension. Yet another way to slow absorption of a particular compound is by administering injectable depot forms comprising microcapsule matrices of the compound trapped within liposomes, microemulsions, or biodegradable polymers such as polylactide-polyglycolide, polyorthoesters or polyanhydrides. Depending on the ratio of drug to polymer and the composition of the polymer, the rate of drug release can be controlled.

Transdermal patches can also provide controlled delivery of the compounds. The rate of absorption can be slowed by using rate controlling membranes or by trapping the compound within a polymer matrix or gel. Conversely, absorption enhancers can be used to increase absorption.

Solid dosage forms for oral administration include capsules, tablets, pills, powders, and granules. In these solid dosage forms, the active compound can optionally comprise diluents such as sucrose, lactose, starch, talc, silicic acid, aluminum hydroxide, calcium silicates, polyamide powder, tableting lubricants, and tableting aids such as magnesium 5 stearate or microcrystalline cellulose. Capsules, tablets and pills can also comprise buffering agents, and tablets and pills can be prepared with enteric coatings or other release-controlling coatings. Powders and sprays can also contain excipients such as talc, silicic acid, aluminum hydroxide, calcium silicate, polyamide powder, or mixtures thereof. Sprays can additionally contain customary propellants such as chlorofluorohydrocarbons or substitutes therefor.

10 Liquid dosage forms for oral administration include emulsions, microemulsions, solutions, suspensions, syrups, and elixirs comprising inert diluents such as water. These compositions can also comprise adjuvants such as wetting, emulsifying, suspending, sweetening, flavoring, and perfuming agents.

15 Topical dosage forms include ointments, pastes, creams, lotions, gels, powders, solutions, sprays, inhalants, and transdermal patches. The compound is mixed under sterile conditions with a carrier and any needed preservatives or buffers. These dosage forms can also include excipients such as animal and vegetable fats, oils, waxes, paraffins, starch, tragacanth, cellulose derivatives, polyethylene glycols, silicones, bentonites, silicic acid, talc and zinc oxide, or mixtures thereof. Suppositories for rectal or vaginal administration can be 20 prepared by mixing the compounds with a suitable non-irritating excipient such as cocoa butter or polyethylene glycol, each of which is solid at ordinary temperature but fluid in the rectum or vagina. Ophthalmic formulations comprising eye drops, eye ointments, powders, and solutions are also contemplated as being within the scope of the present invention.

25 The total daily dose of the compounds administered to a host in single or divided doses can be in amounts from about 0.1 to about 200 mg/kg body weight or preferably from about 0.25 to about 100 mg/kg body weight. Single dose compositions can contain these amounts or submultiples thereof to make up the daily dose.

Preferred compounds of the present invention include, but are not limited to:

30 Compounds of formula (I) wherein R¹ is aminocarbonyl. Most preferred compounds of the present invention include compounds of formula (I) wherein R¹ is -C(O)NHCH₃.

Determination of Biological Activity

Activity assay for human histone deacetylases

35 Compounds of the present invention were tested in one of two assays designed to measure histone deacetylase inhibition. Conditions for each of the two assays are described below.

Assay 1

Nuclear histone deacetylase enzymes were partially purified from human erythroleukemia K562 cells by MonoQ sepharose chromatography (Proceedings of the National Academy of Sciences of the United States of America **1999**, **96**, 4592). The substrates, (³H)-labeled nuclear histones, were prepared from K562 cells by incubation of cells with (³H)-acetic acid in the presence of 3 mM trichostatin A and isolated by dounce homogenization, acid extraction of isolated nuclei, and acetone precipitation (*J. Biol. Chem.* **1990**, **265**, 17174). The standard assay consisted of 3-6 µg of histone deacetylase incubated with 5-10 µg (~10,000 cpm) of labeled-nuclear histones for 1 hour at 37 °C in a 50 mL reaction volume. Inhibitor was added 15 minutes prior to substrate addition. The reaction was terminated by the addition of 1M HCl/0.16M acetic acid (50 mL) and ethyl acetate (500 mL). The mixture was inverted for 30 seconds and the phases were separated by centrifugation (1000 rpm for 2 minutes). An aliquot of the organic phase was removed and counted in a liquid scintillation spectrophotometer. IC₅₀ values were determined by log-logit linear regression of the dose response data.

Assay 2

A white DYNEX Microfluor 2 plate was treated with 70 µL buffer (containing 10 mM Tris HCl, 1 mM MgCl₂, 10 mM CaCl₂ at pH 8.0 containing 2% glycerol and 0.015% Tween-80); 12 µL inhibitor (compound) solution in 10% DMSO/buffer; and 18 µL HDAC solution diluted in buffer (the amount of HDAC is adjusted to cleave approximately 10% of the acetyl-lysine from the peptide substrate in a 30 minute reaction). The plate was mixed and preincubated for 30 minutes at room temperature, treated with 20 µL of a 4.8µM solution of substrate (a histone mimetic sequence containing one acetyl lysine group, prepared as a 0.24 mM DMSO stock solution), and incubated for 30 minutes. Each well was treated with 30 µL of a solution of endoproteinase-Lys-C containing trichostatin-A (endoproteinase-Lys-C was added at a concentration of 10 ng/well and the final concentration of trichostatin-A was 7 µM in 150 µL). The buffer used for the quench was 10 mM Hepes/5 mM EDTA, adjusted to pH 8.0 with NaOH, and contains 2% glycerol and 0.015% Tween-80.

The wells of the plate were read by a fluorescence plate reader (fmax, Molecular Devices) with filters of 544 nm (excitation) and 590 nm (emission). The background fluorescence was determined by addition of trichostatin-A to certain wells before addition of enzyme, and was subtracted from the readings of the other wells. The extent of inhibition of the enzyme by the inhibitors was calculated from the readings of wells containing an inhibitor and those of control (containing no inhibitor). The IC₅₀ was determined by a log/logit analysis of the inhibitor concentration and inhibition data.

The compounds of the present invention were found to inhibit histone deacetylase with inhibitory potencies between 1 nM and 50 μ M. Preferred compounds inhibited histone deacetylase with inhibitory potencies between 1 nM and 1 μ M and most preferred compounds inhibited histone deacetylase with inhibitory potencies between 1 nM and 100 nM. Thus, the compounds of the present are useful for treating diseases in which histone deacetylase plays a role.

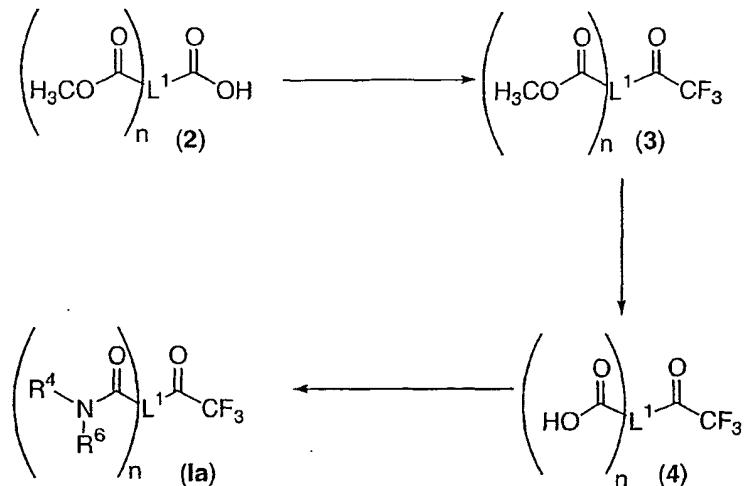
Synthetic Methods

Abbreviations which have been used in the descriptions of the schemes and the examples that follow are: DMAP for 4-dimethylaminopyridine; CDI for 1,1'-carbonyldiimidazole; EDCI for 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride; DCC for 1,3-dicyclohexylcarbodiimide; HOBt for 1-hydroxybenzotriazole hydrate; DMF for N,N-dimethylformamide; NMP for N-methylpyrrolidinone; THF for tetrahydrofuran; MTBE for methyl tert-butyl ether; DMSO for dimethylsulfoxide; OAc for acetate; DME for 1,2-dimethoxyethane; DEAD for diethyl azodicarboxylate; DIAD for diisopropyl azodicarboxylate; LAH for lithium aluminum hydride; NMM for N-methylmorpholine; TBAF for tetrabutylammonium fluoride; DBU for 1,8-diazabicyclo(5.4.0)undec-7-ene; pTsOH for p-toluenesulfonic acid; DBN for 1,5-diazabicyclo(4.3.0)non-5-ene; LDA for lithium diisopropylamide; KHMDS for potassium hexamethyldisilazide; PDC for pyridinium dichromate; NBS for N-bromosuccinimide; TBS for tert-butyldimethylsilyl, and mCPBA for m-chloroperoxybenzoic acid.

The compounds and processes of the present invention will be better understood in connection with the following synthetic schemes which illustrate the methods by which the compounds of the invention may be prepared. Starting materials can be obtained from commercial sources or prepared by well-established literature methods known to those of ordinary skill in the art. The groups n, L¹, L², R¹, R², R³, R⁴, R⁵, and R⁶ are as defined above unless otherwise noted below.

This invention is intended to encompass compounds having formula (I) when prepared by synthetic processes or by metabolic processes. Preparation of the compounds of the invention by metabolic processes include those occurring in the human or animal body (*in vivo*) or processes occurring *in vitro*.

Scheme 1

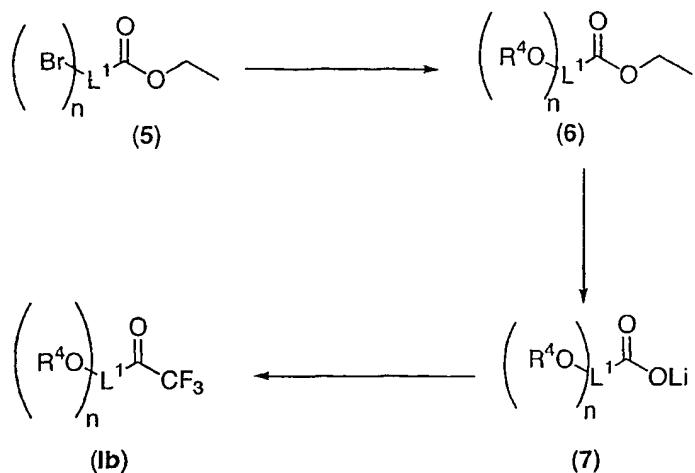


As shown in Scheme 1, compounds of formula (2) can be converted to compounds of formula (3) by treatment with a base and trifluoroacetic anhydride. Examples of bases used in these reactions include sodium hydride, lithium hexamethyldisilazide, pyridine, and mixtures thereof. Representative solvents used in these reactions include dichloromethane, carbon tetrachloride, and chloroform. The reaction is conducted at about -10 °C to about 5 °C and reaction times are typically about 2 to about 24 hours.

Compounds of formula (3) can be converted to compounds of formula (4) by hydrolysis methods known to those of ordinary skill in the art.

Conversion of compounds of formula (4) to compounds of formula (1a) can be accomplished by treatment with an appropriately substituted amine (HNR^4R^6) in the presence of a base and a coupling agent. Examples of bases include NMM, DMAP, and triethylamine. Representative coupling agents include CDI, EDCI, DCC, HOEt, and mixtures thereof. Solvents typically used in these reactions include DMF, NMP, and dioxane. The reaction is conducted at about 20 °C to about 40 °C and reaction times are typically about 12 to about 24 hours.

Scheme 2



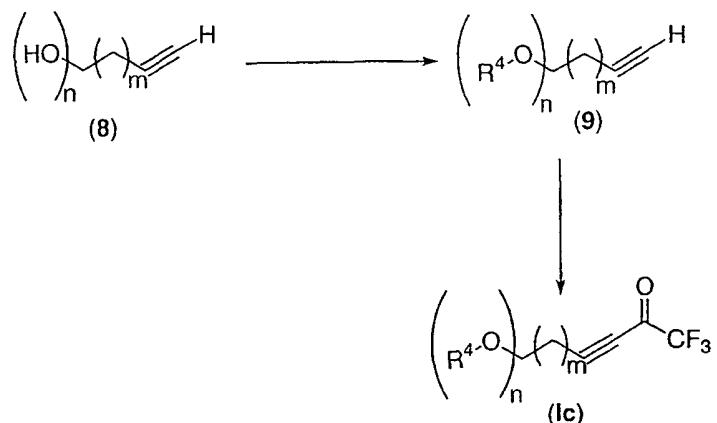
As shown in Scheme 2, compounds of formula (5) can be converted to compounds of formula (6) by treatment with an appropriately substituted alcohol (R^4OH) in the presence of a base. Example of bases include Cs_2CO_3 , K_2CO_3 , and Na_2CO_3 . Representative solvents include DMF, NMP, and dioxane. The reaction is conducted at about 20 °C to about 40 °C and reaction times are typically about 12 to about 24 hours.

Compounds of formula (6) can be converted to compounds of formula (7) by hydrolysis methods known to those of ordinary skill in the art.

Conversion of compounds of formula (7) to compounds of formula (Ib) can be accomplished by treatment with trifluoroacetic anhydride. Solvents commonly used in these reactions include dichloromethane, chloroform, and carbon tetrachloride. The reaction is conducted at about 20 °C to about 40 °C and reaction times are typically about 2 to about 4 hours.

Compounds of formula (Ib) wherein L^1 is alkynylene or alkenylene can be readily converted to compounds of formula (Ib) wherein L^1 is alkenylene or cycloalkylene, respectively, by methods such as cyclopropanation and reduction, well-known to those of ordinary skill in the art.

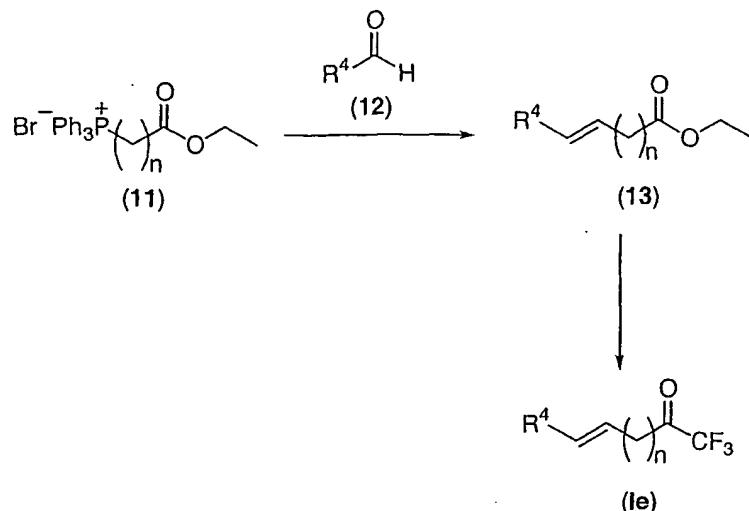
Scheme 3



As shown in Scheme 3, compounds of formula (8) (m is a positive integer between 1 and 7) can be treated with an appropriately substituted alcohol (R⁴OH) in the presence of a trialkylphosphine or triarylphosphine and a diazo compound to provide compounds of formula (9). Representative trialkylphosphines include tributylphosphine and trimethylphosphine; representative triarylphosphines include triphenylphosphine and tri-o-tolylphosphine; and representative diazo compounds include DEAD and DIAD. Solvents commonly used in these reactions include THF, diethyl ether, and methyl tert-butyl ether. The reaction is conducted at about -5 °C to about 30 °C, and typical reaction times are about 12 to about 24 hours.

Compounds of formula (9) can be converted to compounds of formula (Ic) by treatment with base followed by an ester of trifluoroacetic acid. Representative bases include n-butyllithium, tert-butyllithium, and lithium hexamethyldisilazide. Examples of solvents used in these reactions include THF, diethyl ether, and methyl tert-butyl ether. The reaction is conducted at about -78 °C to about 30 °C and typical reaction times are about 18 to about 24 hours.

Scheme 4



As shown in Scheme 4, compounds of formula (11) (n is a positive integer between 1 and 8) can be treated with compounds of formula (12) and base to provide compounds of formula (13). Examples of bases used in these reactions include potassium tert-butoxide and sodium tert-butoxide. Representative solvents include THF, methyl tert-butyl ether, and diethyl ether. The reaction temperature is about -5 °C to about 25 °C and reaction times are typically about 1 to about 3 hours.

Compounds of formula (13) can be converted to compounds of formula (1e) following the procedures described in Scheme 2.

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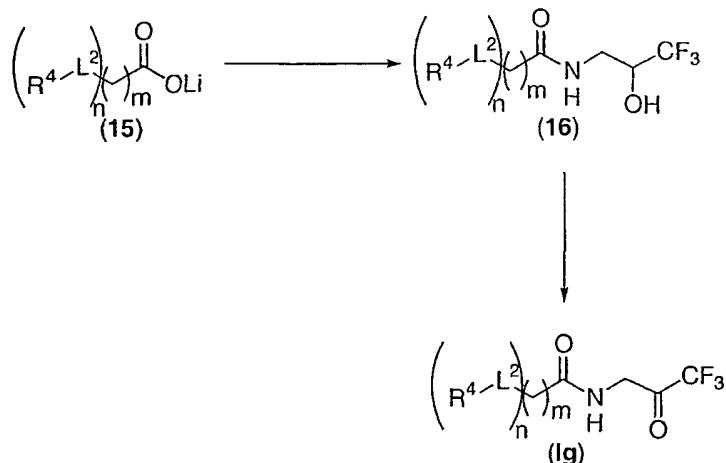
Scheme 5



As shown in Scheme 5, compounds of formula (14) can be converted to compounds of formula (1f) by treatment with oxalyl chloride, followed by treatment with trifluoroacetic anhydride and base. Examples of bases include pyridine, triethylamine, and diisopropylethylamine. Representative solvents include dichloromethane, 1,2-dichloroethane, and carbon tetrachloride. The reaction temperature is about -60 °C to about 25 °C and reaction times are typically about 2 hours to about 4 hours.

20

Scheme 6



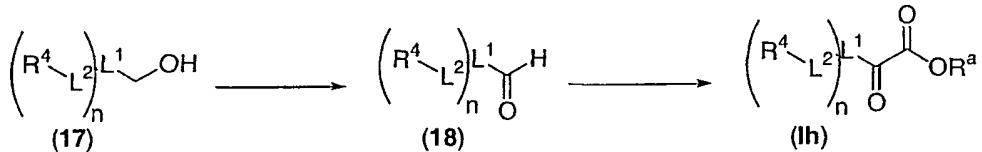
As shown in Scheme 6, compounds of formula (15) (prepared from the corresponding ester according to the procedures described in Scheme 2) can be converted to compounds of formula (16) by treatment with 2-hydroxy-2-(trifluoromethyl)ethylamine (prepared as

5 described in *J. Org. Chem.* 1995, 60, 41) using the conditions described in Scheme 2.

Conversion of compounds of formula (16) to compounds of formula (1g) can be accomplished by oxidation, using a variety of procedures known to those of ordinary skill in the art.

10

Scheme 7



As shown in Scheme 7, compounds of formula (17) can be oxidized to compounds of formula (18) by numerous methods well-known to those of ordinary skill in the art, such as the Swern oxidation and the Dess-Martin oxidation.

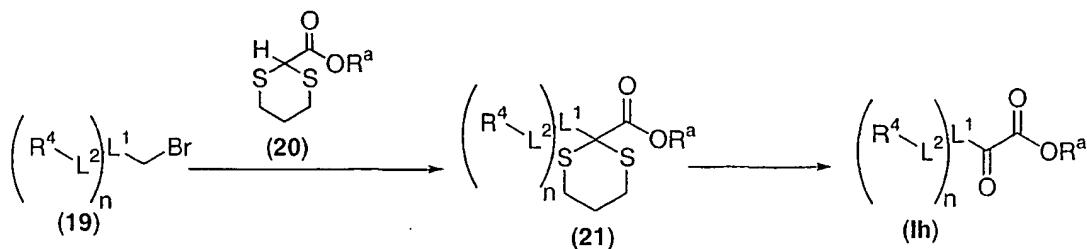
15

Compounds of formula (18) can be converted to compounds of formula (1h) (R^a is alkyl) by treatment with an alkyl ester of (dimethoxyphosphoryl)(tetrahydro-2H-pyran-2-yloxy)acetic acid (which can be prepared following the procedure described in *Tet. Lett.* 1981, 22, 663-666) in the presence of base. Representative bases include DBU, DBN, and DMAP. Examples of solvents used in these reactions include acetonitrile, THF, and diethyl ether. The reaction is conducted at about 0 °C to about 25 °C and reaction times are typically about 1 to about 3 hours.

Compounds of formula (1h) where R^a is alkyl can be intraconverted to compounds of formula (1h) where R^a is hydrogen by the hydrolysis methods shown in Scheme 1.

25

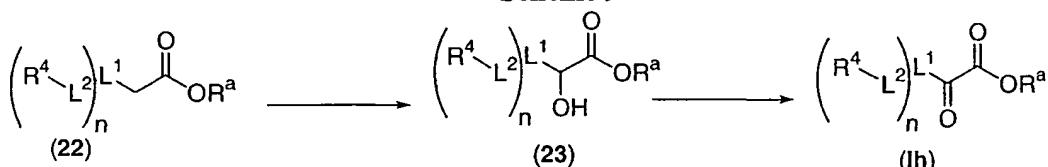
Scheme 8



Scheme 8 shows an alternative synthesis of compounds of formula (Ih). Compounds of formula (19) can be reacted with compounds of formula (20) (R^a is alkyl) in the presence of base to provide compounds of formula (21). Representative bases include NaH, KH, and LiHMDS. Examples of solvents used in these reactions include DMF, THF, and diethyl ether. The reaction is conducted at about -78°C to about 0°C and reaction times are typically about 12 to about 24 hours.

Compounds of formula (21) can be converted to compounds of formula (Ih) by treatment with a variety of deprotection reagents such as NBS, known to those of ordinary skill in the art.

Scheme 9

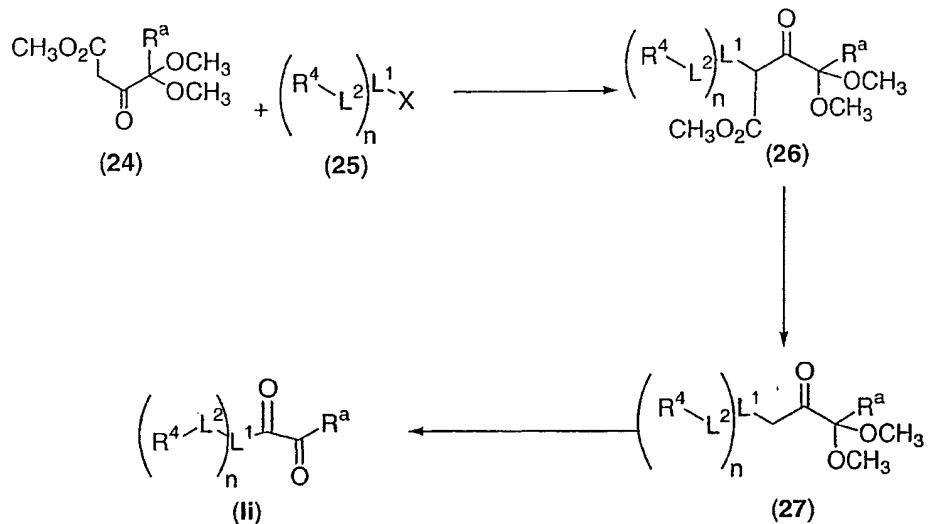


Another route to compounds of formula (Ih) is shown in Scheme 9. Compounds of formula (22) can be converted to compounds of formula (23) by treatment with 2-benzenesulfonyl-3-phenyl-oxaziridine (prepared according to the procedure described in J. Org. Chem. 1992, 47, 1774-1775) in the presence of base. Representative bases include KHMDS, LiHMDS, and LDA. Examples of solvents used in these reactions include THF, MTBE, and diethyl ether. The reaction is conducted at about -78°C to about 0°C and reaction times are typically about 30 minutes to about 2 hours.

Compounds of formula (23) can be converted to compounds of formula (Ih) by treatment with a variety of oxidation reagents, such as PDC, known to those of ordinary skill in the art.

Compounds of formula (Ih) can be reacted with various primary or secondary amines to form the corresponding ketoamides using procedures well-known to those of ordinary skill in the art.

Scheme 10

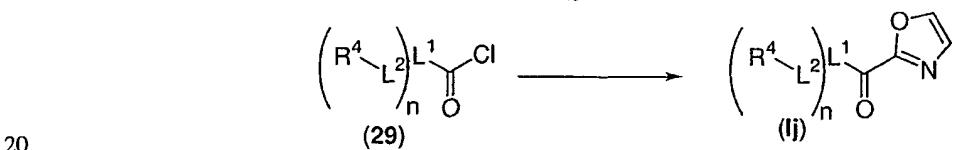


As shown in Scheme 10, compounds of formula (24) (R^a is alkyl) can be treated sequentially with a base and with compounds of formula (25) to provide compounds of formula (26). Representative bases include sodium hydride, potassium hydride, lithium hexamethyldisilazide, and lithium diisopropylamide. Examples of solvents used in these reactions include DMF, THF, MTBE, and diethyl ether. The reaction is typically conducted at about -78°C to about 25°C for about 2 to about 48 hours.

Compounds of formula (26) can be converted to compounds of formula (27) by treatment with a hydrolyzing agent. Representative hydrolyzing agents include sodium hydroxide and lithium hydroxide. Examples of solvents used in these reactions include methanol and ethanol. The reaction is conducted at about 25°C to about 75°C for about 1 to about 6 hours.

Conversion of compounds of formula (27) to compounds of formula (Ii) can be accomplished by treatment with a deprotecting agent such as HCl, trifluoroacetic acid, p-toluenesulfonic acid, or acetic acid. The reaction is conducted at about 0°C to about 35°C for about 1 to about 12 hours.

Scheme 11

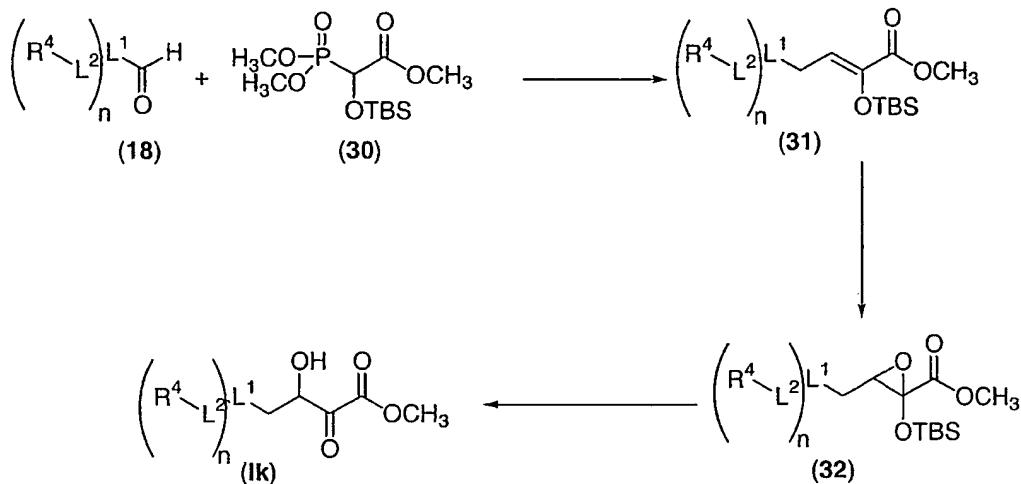


Scheme 11 shows the conversion of compounds of formula (29) to compounds of formula (Ii). Treatment of compounds of formula (29) with a stabilized anion of a heterocycle (generated by deprotonation with a strong base such as n-butyllithium at

-78 °C followed by treatment with zinc chloride) in the presence of stoichiometric copper (such as copper iodide) gives compounds of formula (Ij). Examples of solvents used in these reactions include THF, diethyl ether, and MTBE. The reaction is conducted at about -78 °C to about 0 °C and reaction times are typically about 1 to about 3 hours.

5

Scheme 12

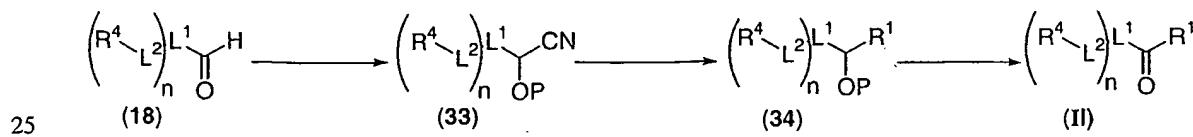


As shown in Scheme 12, compounds of formula (18) (prepared according to the methods described in Scheme 7) can be reacted with compounds of formula (30) in the presence of base and lithium chloride to provide compound of formula (31). Examples of bases include DBU, diisopropylethylamine, and sodium hydride. Representative solvents include THF, MTBE, and dioxane. The reaction is conducted at about 0 to about 23 °C for about 1 to about 16 hours.

Compounds of formula (31) can be reacted with an oxidizing agent to produce compounds of formula (32). Representative oxidizing agents include mCPBA with potassium fluoride, and t-butyl peroxide with n-butyllithium. Examples of solvents include dichloromethane, THF, and chloroform. The reaction is conducted at about 0 to about 23 °C for about 8 to about 16 hours.

Conversion of compounds of formula (32) to compounds of formula (Ik) can be accomplished by treatment with triethylamine trihydrofluoride. Examples of solvents used in this reaction include acetonitrile, tetrahydrofuran, and toluene. The reaction is conducted at about 0 to about 23 °C for about 8 to about 16 hours.

Scheme 13



25

As shown in Scheme 13, compounds of formula (18) can be reacted with KCN to provide compounds of formula (33) (P is H). Examples of solvents used in these reactions include THF, water, and mixtures thereof. The reaction is typically conducted at about 10 °C to about 35 °C for about 12 to about 72 hours.

5 Compounds of formula (33) where P is H can be converted to compounds of formula (33) where P is a hydroxy protecting group can be accomplished by means known to those of ordinary skill in the art.

10 Compounds of formula (33) where P is H can be converted to compounds of formula (34) (where R is dihydrooxazolyl by treatment with acetyl chloride in ethanol and chloroform followed by treatment with ethanolamine and triethylamine in dichloromethane, followed by treatment with p-toluenesulfonic acid in chloroform. Reaction temperatures are typically between 20 °C and 62 °C and reaction times are typically about 3 to about 24 hours.

15 Compounds of formula (33) can be converted to compounds of formula (33) where P is a hydroxy protecting group can be converted to compounds of formula (34) by treatment with sodium azide and ammonium chloride in DMF. Reaction temperatures are typically about 80 °C to about 153 °C for about 1 to about 6 hours.

Compounds of formula (34) where P is a hydroxy protecting group can be converted to compounds of formula (34) where P is hydrogen by methods known to those of ordinary skill in the art.

20 Conversion of compounds of formula (34) to compounds of formula (II) can be accomplished by oxidation using methods known to those of ordinary skill in the art.

25 The present invention will now be described in connection with certain preferred embodiments which are not intended to limit its scope. On the contrary, the present invention covers all alternatives, modifications, and equivalents as can be included within the scope of the claims. Thus, the following examples, which include preferred embodiments, will illustrate the preferred practice of the present invention, it being understood that the examples are for the purposes of illustration of certain preferred embodiments and are presented to provide what is believed to be the most useful and readily understood description of its procedures and conceptual aspects.

30 Compounds of the invention were named by ACD/ChemSketch version 5.0 (developed by Advanced Chemistry Development, Inc., Toronto, ON, Canada) or were given names which appeared to be consistent with ACD nomenclature.

Example 1

35 9,9,9-trifluoro-8-oxo-N-phenylnonanamide

Example 1A

methyl 9,9,9-trifluoro-8-oxononanoate

A suspension of NaH (750 mg, 29.7 mmol) in dichloromethane (150 mL) at 0 °C was treated dropwise with 8-methoxy-8-oxooctanoic acid (5.10 g, 27.1 mmol), stirred until gas evolution ceased, treated with trifluoroacetic anhydride (34.2 g, 163 mmol), stirred for 10 minutes, and treated with pyridine (18.9 g, 225 mmol). The mixture was warmed to room temperature, stirred for 1.5 hours, poured over ice (400 g), and warmed to room temperature. The layers were separated and the aqueous phase was extracted with dichloromethane. The combined extracts were dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 1:1 dichloromethane/hexanes to provide 2.68 g (41%) of the desired product. MS (ESI(-)) m/e 239 (M-H)⁻.

Example 1B9,9,9-trifluoro-8-oxononanoic acid

A solution of Example 1A (1.40 g, 5.8 mmol) in THF (25 mL) at room temperature was treated with 2M LiOH (35 mL, 70 mmol), stirred for 18 hours, and concentrated. The remaining solution was adjusted to pH 2 with 1N HCl and extracted with ethyl acetate. The combined extracts were dried (Na₂SO₄), filtered, and concentrated to provide 1.28 g (98%) of the desired product of sufficient purity for subsequent use. MS (ESI(-)) m/e 225 (M-H)⁻.

20

Example 1C9,9,9-trifluoro-8-oxo-N-phenylnonanamide

A solution of Example 1B (256 mg, 1.1 mmol), aniline (112 mg, 1.2 mmol), HOBT (179 mg, 1.3 mmol) and N-methylmorpholine (221 mg, 2.2 mmol) in DMF (3 mL) at room temperature was treated with EDCI (254 mg, 1.3 mmol), stirred for 18 hours, poured into water (50 mL), and extracted with ethyl acetate. The combined extracts were dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 7:3 hexanes/ethyl acetate to provide 273 mg (82%) of the desired product. MS (ESI(+)) m/e 302 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.83 (s, 1H), 7.58 (d, 2H), 7.35-7.30 (m, 2H), 7.01 (td, 1H), 2.86 (t, 2H), 2.29 (t, 2H), 1.65-1.50 (m, 4H), 1.47-1.23 (m, 4H); Anal. Calcd for C₁₅H₁₈F₃NO₂: C, 59.79; H, 6.02; N, 4.65. Found: C, 59.62; H, 5.91; N, 4.51.

Example 28-((1,1'-biphenyl)-4-yloxy)-1,1,1-trifluoro-2-octanone

35

Example 2Aethyl 7-((1,1'-biphenyl)-4-yloxy)heptanoate

A mixture of ethyl 7-bromoheptanoate (53.1 g, 15.4 mmol), (1,1'-biphenyl)-4-ol (2.61 g, 15.3 mmol), and Cs₂CO₃ (5.49 g, 16.9 mmol) in DMF (50 mL) at room temperature was stirred for 18 hours, poured into ice water (400 mL), and filtered to provide 4.87 g (98%) of the desired product. MS (ESI(+)) m/e 327 (M+H)⁺.

5

Example 2B

lithium 7-((1,1'-biphenyl)-4-yloxy)heptanoate

A solution of Example 2A (4.86 g, 14.9 mmol) in THF (15 mL) at room temperature was treated with 2M LiOH (25 mL, 50 mmol), heated to 80 °C for 2 hours, cooled to room 10 temperature, filtered, and dried to provide 4.30g (95%) of the desired product. MS (ESI(-)) m/e 297 (M-Li)⁻.

Example 2C

8-((1,1'-biphenyl)-4-yloxy)-1,1,1-trifluoro-2-octanone

A solution of trifluoroacetic anhydride (2.14 g, 10.2 mmol) in dichloromethane (16 mL) at room temperature was treated with Example 2B (502 mg, 1.66 mmol), and pyridine (1.12 g, 13.4 mmol), stirred for 3 hours, and quenched with water (5 mL). The mixture was stirred for 10 minutes, poured into water (75 mL), and extracted with dichloromethane. The combined extracts were dried (Na₂SO₄), filtered, and concentrated. The concentrate was 20 purified by flash column chromatography on silica gel with dichloromethane to provide 235 mg (40%) of the desired product. MS (ESI(-)) m/e 349 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 7.62-7.56 (m, 4H), 7.45-7.40 (m, 2H), 7.37-7.33 (m, 1H), 7.00 (d, 2H), 4.00 (t, 2H), 2.89 (t, 2H), 1.75-1.68 (m, 2H), 1.65-1.55 (m, 2H), 1.47-1.36 (m, 4H); Anal. Calcd for C₂₀H₂₁F₃O₂: C, 68.56; H, 6.04. Found: C, 68.35; H, 6.10.

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Example 3

4'-((8,8,8-trifluoro-7-oxooctyl)oxy)(1,1'-biphenyl)-4-carbonitrile

The desired product was prepared by substituting 4'-hydroxy(1,1'-biphenyl)-4-carbonitrile for (1,1'-biphenyl)-4-ol in Example 2. MS (ESI(-)) m/e 374 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 7.87 (d, 2H), 7.83 (d, 2H), 7.70 (d, 2H), 7.05 (d, 2H), 4.03 (t, 2H), 2.89 (t, 2H), 1.76-1.69 (m, 2H), 1.62-1.57 (m, 2H), 1.49-1.36 (m, 4H); Anal. Calcd for C₂₁H₂₀F₃NO₂: C, 67.19; H, 5.37; N, 3.73. Found: C, 67.24; H, 5.29; N, 3.58.

Example 4

9-((1,1'-biphenyl)-4-yloxy)-1,1,1-trifluoro-2-nanone

The desired product was prepared by substituting ethyl 8-bromoocanoate for ethyl 7-bromoheptanoate in Example 2. MS (ESI(-)) m/e 363 (M-H)⁻; ¹H NMR (300 MHz, DMSO-

d₆) δ 7.62-7.55 (m, 4H), 7.45-7.39 (m, 2H), 7.33-7.27 (m, 1H), 7.01 (d, 2H), 4.00 (t, 2H), 2.88 (t, 2H), 1.75-1.68 (m, 2H), 1.60-1.55 (m, 2H), 1.43-1.32 (m, 6H); Anal. Calcd for C₂₁H₂₃F₃O₂: C, 69.22; H, 6.36. Found: C, 69.12; H, 6.28.

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Example 57-((1,1'-biphenyl)-4-yloxy)-1,1,1-trifluoro-2-heptanone

The desired product was prepared by substituting ethyl 6-bromohexanoate for ethyl 7-bromoheptanoate in Example 2. MS (ESI(-)) m/e 335 (M-H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 7.62-7.57 (m, 4H), 7.45-7.40 (m, 2H), 7.33-7.27 (m, 1H), 7.01 (d, 2H), 4.01 (t, 2H), 2.92 (t, 2H), 1.78-1.71 (m, 2H), 1.71-1.62 (m, 2H), 1.51-1.43 (m, 2H); Anal. Calcd for C₁₉H₁₉F₃O₂: C, 67.85; H, 5.69. Found: C, 67.82; H, 5.69.

Example 69,9,9-trifluoro-8-oxo-N-(4-pyridinyl)nonanamide

A mixture of Example 1B (50 mg, 0.22 mmol), HOBr (30 mg, 0.22 mmol), carbodiimide PS resin (720 mg), and 4-aminopyridine (25 mg, 0.27 mmol) in DMF (5 mL) at room temperature was agitated in a Quest 210 parallel synthesizer for 18 hours, treated with trisamine PS resin (220 mg), and agitated for 2 hours. The solution was decanted, the resin was rinsed with dichloromethane, and the combined solutions were concentrated. The concentrate was purified by preparative HPLC with a gradient system of 0 to 95% over 10 min of CH₃CN (containing 0.1%TFA) in water to provide the desired product. MS (ESI(+)) m/e 303 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 11.11 (s, 1H), 8.64 (d, 2H), 7.96 (d, 2H), 2.87 (t, 2H), 2.46 (t, 2H), 1.63-1.55 (m, 4H), 1.34-1.29 (m, 4H); Anal. Calcd for C₁₆H₂₀F₃NO₂·CF₃CO₂H·0.1H₂O: C, 45.96; H, 4.39; N, 6.70. Found: C, 45.60; H, 4.30; N, 6.70.

Example 7N-benzyl-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting benzylamine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 316 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.32-8.25 (m, 1H), 7.33-7.28 (m, 2H), 7.26-7.20 (m, 3H), 4.25 (d, 2H), 2.85 (t, 2H), 2.13 (t, 2H), 1.64-1.46 (m, 4H), 1.33-1.28 (m, 4H); Anal. Calcd for C₁₆H₂₀F₃NO₂·0.75H₂O: C, 58.44; H, 6.59; N, 4.26. Found: C, 58.18; H, 6.45; N, 4.05.

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Example 89,9,9-trifluoro-8-oxo-N-(3-pyridinylmethyl)nonanamide

The desired product was prepared by substituting 3-pyridinylmethanamine for 4-

aminopyridine in Example 6. MS (ESI(+)) m/e 317 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.61-8.58 (m, 2H), 8.44-8.40 (m, 1H), 7.97-7.94 (m, 1H), 7.62 (dd, 1H), 4.34 (d, 2H), 2.85 (t, 2H), 2.14 (t, 2H), 1.62-1.51 (m, 4H), 1.28-1.22 (m, 4H).

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Example 9

9,9,9-trifluoro-8-oxo-N-(2-phenylethyl)nonanamide

The desired product was prepared by substituting 2-phenylethanamine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 330 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 7.84 (m, 1H), 7.31-7.26 (m, 2H), 7.21-7.18 (m, 3H), 3.25 (q, 2H), 2.85 (t, 2H), 2.69 (t, 2H), 2.02 (t, 2H), 1.58-1.43 (m, 4H), 1.30-1.18 (m, 4H).

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Example 10

9,9,9-trifluoro-N-(4-methoxyphenyl)-8-oxononanamide

The desired product was prepared by substituting 4-methoxyaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 332 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.68 (s, 1H), 7.47 (d, 2H), 6.85 (d, 2H), 3.71 (s, 3H), 2.88-2.84 (m, 2H), 2.28-2.23 (m, 2H), 1.64-1.52 (m, 4H), 1.34-1.25 (m, 4H); Anal. Calcd for C₁₆H₂₀F₃NO₃·0.7H₂O: C, 55.87; H, 6.27; N, 4.07. Found: C, 55.64; H, 6.13; N, 3.88.

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Example 11

9,9,9-trifluoro-N-(3-methoxyphenyl)-8-oxononanamide

The desired product was prepared by substituting 3-methoxyaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 332 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.84 (s, 1H), 7.30 (br s, 1H), 7.21-7.09 (m, 2H), 6.61-6.59 (m, 1H), 3.71 (s, 3H), 2.87 (t, 2H), 2.28 (t, 2H), 1.62-1.51 (m, 4H), 1.36-1.25 (m, 4H); Anal. Calcd for: C₁₆H₂₀F₃NO₃·0.6H₂O: C, 56.17; H, 6.25; N, 4.09. Found: C, 55.81; H, 6.04; N, 3.91.

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Example 12

9,9,9-trifluoro-N-(2-methoxyphenyl)-8-oxononanamide

The desired product was prepared by substituting 2-methoxyaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 332 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.99 (s, 1H), 7.91 (d, 1H), 7.08-7.00 (m, 2H), 6.91-6.85 (m, 1H), 3.82 (s, 3H), 2.87 (t, 2H), 2.36 (t, 2H), 1.64-1.51 (m, 4H), 1.34-1.26 (m, 4H).

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Example 13

9,9,9-trifluoro-8-oxo-N-(3-phenylpropyl)nonanamide

The desired product was prepared by substituting 3-phenyl-1-propanamine for 4-

aminopyridine in Example 6. MS (ESI(+)) m/e 344 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 7.78 (br t, 1H), 7.30-7.25 (m, 2H), 7.20-7.14 (m, 3H), 3.03 (dd, 2H), 2.85 (t, 2H), 2.58-2.53 (m, 2H), 2.05 (t, 2H), 1.72-1.43 (m, 6H), 1.31-1.21 (m, 4H); Anal. Calcd for C₁₈H₂₄F₃NO₂·H₂O: C, 59.82; H, 7.25; N, 3.88. Found: C, 59.42; H, 6.94; N, 3.80.

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Example 14

N-(4-(dimethylamino)phenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 4-dimethylaminoaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 345 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.68 (br s, 1H), 7.47 (br d, 2H), 6.95-6.83 (br s, 1H), 6.64-6.54 (br s, 1H), 2.92 (br s, 6H), 2.86 (t, 2H), 2.25 (t, 2H), 1.65-1.52 (m, 4H), 1.33-1.25 (m, 4H).

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Example 15

N-(1,3-benzodioxol-5-yl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 1,3-benzodioxol-5-amine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 346 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.76 (br s, 1H), 7.31-7.30 (br s, 1H), 6.95-6.92 (m, 1H), 6.82 (d, 1H), 5.96 (s, 2H), 2.87 (t, 2H), 2.25 (t, 2H), 1.62-1.52 (m, 4H), 1.35-1.26 (m, 4H); Anal. Calcd for C₁₆H₁₈F₃NO₄: C, 55.65; H, 5.25; N, 4.06. Found: C, 55.18; H, 5.00; N, 4.51.

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Example 16

9-(1,4-dioxa-8-azaspiro(4.5)dec-8-yl)-1,1,1-trifluoro-9-oxo-2,2-nonanedio

The desired product was prepared by substituting 1,4-dioxa-8-azaspiro(4.5)decane for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 352 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 6.65-6.48 (br s, 2H), 3.89 (s, 4H), 3.51-3.44 (m, 4H), 2.30 (t, 2H), 1.62-1.56 (m, 4H), 1.55-1.34 (m, 6H), 1.32-1.21 (m, 4H).

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Example 17

N-(1,1'-biphenyl)-4-yl-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting (1,1'-biphenyl)-4-amine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 378 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.96 (s, 1H), 7.70-7.59 (m, 6H), 7.46-7.42 (m, 2H), 7.35-7.30 (m, 1H), 2.88 (t, 2H), 2.35-2.30 (m, 2H), 1.67-1.52 (m, 4H), 1.38-1.29 (m, 4H); Anal. Calcd for C₂₁H₂₂F₃NO₂·0.2H₂O: C, 66.20; H, 5.93; N, 3.68. Found: C, 66.06; H, 5.89; N, 3.67.

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Example 18

N-(1,1'-biphenyl)-3-yl-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting (1,1'-biphenyl)-3-amine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 378 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.95 (s, 1H), 7.92 (s, 1H), 7.61-7.56 (m, 3H), 7.50-7.45 (m, 2H), 7.40-7.30 (m, 3H), 2.87 (t, 2H), 2.32 (t, 2H), 1.68-1.52 (m, 4H), 1.39-1.27 (m, 4H).

5

Example 19

N-(1,1'-biphenyl)-2-yl-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting (1,1'-biphenyl)-2-amine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 378 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.18 (s, 1H), 7.42-7.30 (m, 9H), 2.85 (t, 2H), 2.13-2.11 (m, 2H), 1.63-1.33 (m, 4H), 1.28-1.14 (m, 4H).

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Example 20

N-(4-cyclohexylphenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 4-cyclohexylaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 384 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.74 (s, 1H), 7.46 (d, 2H), 7.11 (d, 2H), 2.86 (t, 2H), 2.49-2.38 (m, 1H), 2.26 (t, 2H), 1.82-1.65 (m, 5H), 1.64-1.51 (m, 4H), 1.41-1.25 (m, 9H). Anal. Calcd for C₂₁H₂₈F₃NO₂·0.3H₂O: C, 64.86; H, 7.41; N, 3.60. Found: C, 64.69; H, 7.16; N, 3.36.

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Example 21

9,9,9-trifluoro-8-oxo-N-(4-(1-piperidinyl)phenyl)nonanamide4

The desired product was prepared by substituting 4-(1-piperidinyl)aniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 385 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 10.14-9.84 (br s, 1H), 7.68-7.55 (br s, 2H), 6.66-6.56 (br s, 2H), 2.87 (t, 2H), 2.51 (br s, 4H), 2.33-2.26 (m, 2H), 1.84-1.71 (m, 4H), 1.64-1.52 (m, 5H), 1.49-1.37 (m, 1H), 1.33-1.25 (m, 4H).

25

Example 22

9,9,9-trifluoro-N-(4-(4-morpholinyl)phenyl)-8-oxononanamide

The desired product was prepared by substituting 4-(4-morpholinyl)aniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 387 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.66 (s, 1H), 7.45 (d, 2H), 6.19 (d, 2H), 3.75-3.72 (m, 4H), 3.17-3.04 (m, 4H), 2.86 (t, 2H), 2.25 (t, 2H), 1.63-1.50 (m, 4H), 1.36-1.24 (m, 4H).

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Example 23

N-((1S)-1-benzyl-2-(methylamino)-2-oxoethyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting (2S)-2-amino-N-methyl-3-phenylpropanamide for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 387 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.03-7.98 (m, 1H), 7.92-7.84 (m, 1H), 7.31-7.13 (m, 5H), 4.49-4.38 (m, 1H), 2.99-2.93 (m, 1H), 2.82 (t, 2H), 2.71 (dd, 1H), 2.56 (d, 3H), 2.02 (t, 2H), 1.55-1.44 (m, 2H), 1.36-1.24 (m, 2H), 1.24-1.11 (m, 2H), 1.11-1.06 (m, 2H); Anal. Calcd for C₁₉H₂₅F₃N₂O₃·0.1H₂O: C, 58.78; H, 6.54; N, 7.22. Found: C, 58.40; H, 6.57; N, 7.24.

Example 24

N-benzhydryl-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting benzhydrylamine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 392 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.70 (d, 1H), 7.35-7.20 (m, 10H), 6.11 (d, 1H), 2.83 (t, 2H), 2.20 (t, 2H), 1.58-1.49 (m, 4H), 1.32-1.21 (m, 4H); Anal. Calcd for C₂₂H₂₄F₃NO₂: C, 67.51; H, 6.18; N, 3.58. Found: C, 67.22; H, 6.13; N, 3.57.

15

Example 25

9,9,9-trifluoro-8-oxo-N-(3-pyridinyl)nonanamide

The desired product was prepared by substituting 3-aminopyridine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 321 ($M+H_2O+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 10.13 (br s, 1H), 8.77-8.76 (m, 1H), 8.27-8.25 (m, 1H), 8.08-8.04 (m, 1H), 7.40-7.36 (m, 1H), 2.87 (t, 2H), 2.34 (t, 2H), 1.65-1.56 (m, 4H), 1.36-1.29 (m, 4H).

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Example 26

N-cyclohexyl-9,9,9-trifluoro-8,8-dihydroxynonanamide

The desired product was prepared by substituting cyclohexylamine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 308 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 7.61 (d, 1H), 6.60 (s, 2H), 3.56-3.43 (m, 1H), 2.01 (t, 2H), 1.74-1.36 (m, 12H), 1.34-1.03 (m, 8H); Anal. Calcd for C₁₅H₂₄F₃NO₂·0.9H₂O: C, 55.68; H, 8.04; N, 4.33. Found: C, 55.60; H, 8.19; N, 4.41.

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Example 27

9,9,9-trifluoro-N-(4-hydroxyphenyl)-8-oxononanamide

The desired product was prepared by substituting 4-hydroxyaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 318 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.56 (s, 1H), 9.11 (s, 1H), 7.34 (d, 2H), 6.64 (d, 2H), 2.86 (t, 2H), 2.23 (t, 2H), 1.62-1.52 (m, 4H), 1.34-1.26 (m, 4H); Anal. Calcd for C₁₅H₁₈F₃NO₃: C, 56.78; H, 5.72; N, 4.41. Found: C, 56.73; H, 5.94; N, 4.38.

Example 289,9,9-trifluoro-N-(4-fluorophenyl)-8-oxononanamide

5 The desired product was prepared by substituting 4-fluoroaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 320 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.91 (s, 1H), 7.59 (dd, 2H), 7.12 (t, 2H), 2.87 (t, 2H), 2.28 (t, 2H), 1.65-1.52 (m, 4H), 1.36-1.26 (m, 4H); Anal. Calcd for C₁₅H₁₇F₄NO₂: C, 56.43; H, 5.37; N, 4.39. Found: C, 56.73; H, 5.94; N, 4.38.

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Example 29N-(3-cyanophenyl)-9,9,9-trifluoro-8-oxononanamide

15 The desired product was prepared by substituting 3-cyanoaniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 325 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 10.23 (s, 1H), 8.11-8.09 (m, 1H), 7.78 (dt, 1H), 7.54-7.47 (m, 2H), 2.87 (t, 2H), 2.33 (t, 2H), 1.66-1.53 (m, 4H), 1.34-1.27 (m, 4H); Anal. Calcd for C₁₆H₁₇F₃N₂O₂·0.5H₂O: C, 58.89; H, 5.25; N, 8.58. Found: C, 57.64; H, 5.29; N, 7.91.

Example 30N-(4-chlorophenyl)-9,9,9-trifluoro-8-oxononanamide

20 The desired product was prepared by substituting 4-chloroaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 336 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.98 (s, 1H), 7.61 (d, 2H), 7.33 (d, 2H), 2.89-2.84 (m, 2H), 2.30 (t, 2H), 1.64-1.53 (m, 4H), 1.35-1.24 (m, 4H); Anal. Calcd for C₁₅H₁₇ClF₃NO₂·0.9H₂O: C, 51.19; H, 5.38; N, 3.98. Found: C, 51.07; H, 5.44; N, 3.99.

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Example 31N-(4-acetylphenyl)-9,9,9-trifluoro-8-oxononanamide

30 The desired product was prepared by substituting 1-(4-aminophenyl)ethanone for 4-aminopyridine in Example 6. MS (APCI(+)) 344 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 10.21 (s, 1H), 7.91 (d, 2H), 7.71 (d, 2H), 2.87 (t, 2H), 2.50 (s, 3H), 2.35 (t, 2H), 1.63-1.54 (m, 4H), 1.34-1.28 (m, 4H).

Example 32N-(2-adamantyl)-9,9,9-trifluoro-8-oxononanamide

35 The desired product was prepared by substituting 2-adamantanamine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 360 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 7.60 (d, 1H), 3.82 (d, 1H), 2.85 (t, 2H), 2.13 (d, 2H), 1.99-1.94 (m, 2H), 1.84-1.67 (m,

10H), 1.59-1.43 (m, 6H), 1.34-1.23 (m, 4H); Anal. Calcd for $C_{19}H_{28}F_3NO_2 \cdot 0.1CF_3COOH$: C, 62.19; H, 7.64; N, 3.78. Found: C, 62.64; H, 7.01; N, 3.44.

Example 33

5 9,9,9-trifluoro-8-oxo-N-(4-(trifluoromethyl)phenyl)nonanamide

The desired product was prepared by substituting 4-trifluoromethylaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 370 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 10.23 (s, 1H), 7.79 (d, 2H), 7.65 (d, 2H), 2.87 (t, 2H), 2.34 (t, 2H), 1.64-1.57 (m, 4H), 1.35-1.29 (m, 4H).

10

Example 34

N-(3,4-dichlorophenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 3,4-dichloroaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 370 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 10.15 (s, 1H), 7.99 (d, 1H), 7.54 (d, 1H), 7.48-7.45 (dd, 1H), 2.86 (t, 2H), 2.31 (t, 2H), 1.64-1.52 (m, 4H), 1.34-1.28 (m, 4H).

15

Example 35

N-(4-bromophenyl)-9,9,9-trifluoro-8-oxononanamide

20 The desired product was prepared by substituting 4-bromoaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 380 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 10.00 (s, 1H), 7.56 (d, 2H), 7.46 (d, 2H), 2.87 (t, 2H), 2.30 (t, 2H), 1.64-1.53 (m, 4H), 1.34-1.26 (m, 4H); Anal. Calcd for $C_{15}H_{17}BrF_3NO_2 \cdot 0.9H_2O$: C, 45.45; H, 4.78; N, 3.53. Found: C, 45.37; H, 4.54; N, 3.38.

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Example 36

N-(4-benzylphenyl)-9,9,9-trifluoro-8-oxononanamide

30 The desired product was prepared by substituting 4-benzylaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 392 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.77 (s, 1H), 7.48 (d, 2H), 7.30-7.25 (m, 2H), 7.21-7.17 (m, 3H), 7.12 (d, 2H), 3.87 (s, 2H), 2.87 (t, 2H), 2.26 (t, 2H), 1.63-1.50 (m, 4H), 1.34-1.26 (m, 4H); Anal. Calcd for $C_{22}H_{24}F_3NO_2$: C, 67.51; H, 6.18; N, 3.58. Found: C, 67.36; H, 5.90; N, 3.44.

35

Example 37

9,9,9-trifluoro-8-oxo-N-(4-phenoxyphenyl)nonanamide

The desired product was prepared by substituting 4-phenoxyaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 394 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-

5 d_6) δ 9.87 (s, 1H), 7.59 (d, 2H), 7.35 (dd, 2H), 7.12-7.06 (m, 1H), 6.99-6.59 (m, 4H), 2.87 (t, 2H), 2.29 (t, 2H), 1.66-1.52 (m, 4H), 1.48-1.26 (m, 4H); Anal. Calcd for $C_{21}H_{22}F_3NO_3$: C, 64.11; H, 5.64; N, 3.56. Found: C, 64.01; H, 5.61; N, 3.51.

Example 38

9,9,9-trifluoro-8,8-dihydroxy-N-(9-oxo-9H-fluoren-2-yl)nonanamide

10 The desired product was prepared by substituting 2-amino-9H-fluoren-9-one for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 404 ($M+H$)⁺; 1H NMR (300 MHz, DMSO- d_6) δ 10.14 (s, 1H), 7.95 (br s, 1H), 7.71-7.68 (m, 3H), 7.60-7.55 (m, 2H), 7.43-7.28 (m, 1H), 6.61 (s, 2H), 2.33 (t, 2H), 1.65-1.59 (m, 4H), 1.49-1.39 (m, 2H), 1.36-1.28 (m, 4H); Anal. Calcd for $C_{22}H_{20}F_3NO_3 \cdot 0.7H_2O$: C, 63.52; H, 5.18; N, 3.37. Found: C, 63.24; H, 4.77; N, 3.28.

Example 39

N-(4-(benzyloxy)phenyl)-9,9,9-trifluoro-8-oxononanamide

15 The desired product was prepared by substituting 4-(benzyloxy)aniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 408 ($M+H$)⁺; 1H NMR (300 MHz, DMSO- d_6) δ 9.70 (s, 1H), 7.49-7.32 (m, 7H), 6.93 (d, 2H), 5.05 (s, 2H), 2.86 (t, 2H), 2.25 (t, 2H), 1.63-1.52 (m, 4H), 1.34-1.28 (m, 4H).

20

Example 40

9,9,9-trifluoro-N-(3-methoxypropyl)-8-oxononanamide

25 The desired product was prepared by substituting 3-methoxy-1-propanamine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 298 ($M+H$)⁺.

25

Example 41

9,9,9-trifluoro-N-isopentyl-8-oxononanamide

30 The desired product was prepared by substituting 3-methyl-1-butanamine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 296 ($M+H$)⁺; 1H NMR (300 MHz, DMSO- d_6) δ 7.68 (br m, 1H), 3.07-3.00 (m, 2H), 2.85 (t, 2H), 2.02 (t, 2H), 1.62-1.36 (m, 5H), 1.30-1.20 (m, 6H), 0.85 (d, 6H).

Example 42

N-(4'-cyano(1,1'-biphenyl)-3-yl)-9,9,9-trifluoro-8-oxononanamide

35 The desired product was prepared by substituting 3'-amino(1,1'-biphenyl)-4-carbonitrile for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 403 ($M+H$)⁺; 1H NMR (300 MHz, DMSO- d_6) δ 10.02 (s, 1H), 8.01 (s, 1H), 7.94 (d, 2H), 7.81 (d, 2H), 7.61 (app d, 1H),

7.46-7.37 (m, 2H), 2.87 (t, 2H), 2.33 (t, 2H), 1.60-1.55 (m, 4H), 1.34 (m, 4H); Anal. Calcd for $C_{22}H_{21}F_3N_2O_2$: C, 65.66; H, 5.26; N, 6.96. Found: C, 65.50; H, 5.37; N, 7.04.

Example 43

5

N-(3-(benzyloxy)phenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 3-(benzyloxy)aniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 408 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.73 (s, 1H), 7.48-7.26 (m, 6H), 7.21-7.07 (m, 2H), 6.70-6.65 (m, 1H), 5.06 (s, 2H), 2.86 (t, 2H), 2.28 (t, 2H), 1.59-1.55 (m, 4H), 1.33-1.28 (m, 4H); Anal. Calcd for 10 $C_{22}H_{24}F_3NO_3$: C, 64.86; H, 5.94; N, 3.44. Found: C, 65.06; H, 5.95; N, 3.53.

Example 44

9,9,9-trifluoro-8-oxo-N-(3-phenoxyphenyl)nonanamide

The desired product was prepared by substituting 3-phenoxyaniline for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 394 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) 15 δ 9.92 (s, 1H), 7.42-7.27 (m, 5H), 7.15 (t, 1H), 7.02 (d, 2H), 6.67 (dt, 1H), 2.86 (t, 2H), 2.26 (t, 2H), 1.62-1.50 (m, 4H), 1.34-1.24 (m, 4H); Anal. Calcd for $C_{21}H_{22}F_3NO_3 \cdot 0.2H_2O$: C, 63.53; H, 5.69; N, 3.53. Found: C, 63.42; H, 5.62; N, 3.33.

20

Example 45

N-(3-benzoylphenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting (3-aminophenyl)(phenyl)methanone for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 404 ($M-H$)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 10.10 (s, 1H), 8.01 (s, 1H), 7.90 (dt, 1H), 7.75-7.66 (m, 3H), 7.60-7.54 (m, 2H), 25 7.48 (t, 1H), 7.39 (d, 1H), 2.86 (t, 2H), 2.31 (t, 2H), 1.63-1.52 (m, 4H), 1.34-1.25 (m, 4H); Anal. Calcd for $C_{22}H_{22}F_3NO_3 \cdot 0.2H_2O$: C, 64.60; H, 5.52; N, 3.42. Found: C, 64.46; H, 5.34; N, 3.47.

Example 46

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9,9,9-trifluoro-8-oxo-N-(4-phenyl-1,3-thiazol-2-yl)nonanamide

The desired product was prepared by substituting 4-phenyl-1,3-thiazol-2-amine for 4-aminopyridine in Example 6. MS (ESI(+)) m/e 385 ($M+H$)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 12.21 (s, 1H), 7.89 (d, 2H), 7.59 (s, 1H), 7.45-7.40 (m, 2H), 7.35-7.29 (m, 1H), 2.87 (t, 2H), 2.47 (t, 2H), 1.64-1.54 (m, 4H), 1.35-1.29 (m, 4H); Anal. Calcd for $C_{18}H_{19}F_3N_2O_2S$: C, 56.24; H, 4.98; N, 7.29. Found: C, 55.99; H, 4.94; N, 6.96.

Example 47

8-(3-bromophenoxy)-1,1,1-trifluoro-2-octanone

The desired product was prepared by substituting 3-bromophenol for (1,1'-biphenyl)-4-ol in Example 2. MS (ESI(-)) m/e 351 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 7.23 (t, 1H), 7.13-7.08 (m, 2H), 6.96-6.92 (m, 1H), 3.97 (t, 2H), 2.88 (t, 2H), 1.74-1.65 (m, 2H), 1.63-1.54 (m, 2H), 1.46-1.29 (m, 4H); Anal. Calcd for C₁₄H₁₆BrF₃O₂·0.2H₂O: C, 47.13; H, 4.63. Found: C, 46.75; H, 4.75.

Example 481,1,1-trifluoro-8-(3-(4-pyridinyl)phenoxy)-2-octanone

The desired product was prepared by substituting 3-(4-pyridinyl)phenol for (1,1'-biphenyl)-4-ol in Example 2. MS (ESI(+)) m/e 352 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.62 (d, 2H), 7.71 (d, 2H), 7.42 (t, 1H), 7.36-7.31 (m, 2H), 7.05-7.02 (m, 1H), 4.06 (t, 2H), 2.89 (t, 2H), 1.77-1.60 (m, 2H), 1.68-1.58 (m, 2H), 1.49-1.30 (m, 4H); Anal. Calcd for C₁₉H₂₀F₃NO₂·HCl·2.3H₂O: C, 53.16; H, 6.01; N, 3.26. Found: C, 52.80; H, 5.99; N, 3.03.

15

Example 498-(4-bromophenoxy)-1,1,1-trifluoro-2-octanone

The desired product was prepared by substituting 4-bromophenol for (1,1'-biphenyl)-4-ol in Example 2. MS (ESI(-)) m/e 351 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 7.42 (d, 2H), 6.89 (d, 2H), 3.94 (t, 2H), 2.87 (t, 2H), 1.72-1.65 (m, 2H), 1.62-1.54 (m, 2H), 1.46-1.30 (m, 4H); Anal. Calcd for C₁₄H₁₆BrF₃O₂: C, 47.61; H, 4.57. Found: C, 47.88; H, 4.39.

Example 501,1,1-trifluoro-8-(4-phenoxyphenoxy)-2-octanone

The desired product was prepared by substituting 4-phenoxyphenol for (1,1'-biphenyl)-4-ol in Example 2. MS (ESI(-)) m/e 365 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 7.36-7.32 (m, 2H), 7.09-7.03 (m, 1H), 6.99-6.88 (m, 6H), 3.94 (t, 2H), 2.88 (t, 2H), 1.73-1.64 (m, 2H), 1.61-1.54 (m, 2H), 1.40-1.35 (m, 4H); Anal. Calcd for C₂₀H₂₁F₃O₃: C, 65.57; H, 5.78. Found: C, 65.26; H, 5.64.

30

Example 518-((1,1'-biphenyl)-3-yloxy)-1,1,1-trifluoro-2-octanone

The desired product was prepared by substituting (1,1'-biphenyl)-3-ol for (1,1'-biphenyl)-4-ol in Example 2. MS (ESI(-)) m/e 349 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 7.66 (d, 2H), 7.48-7.45 (m, 2H), 7.39-7.33 (m, 2H), 7.22-7.16 (m, 1H), 7.17-7.16 (m, 1H), 6.94-6.91 (m, 1H), 4.05-4.02 (m, 2H), 2.89 (t, 2H), 1.74 (m, 2H), 1.62-1.58 (m, 2H), 1.50-1.33 (m, 4H); Anal. Calcd for C₂₀H₂₁F₃O₂: C, 68.56; H, 6.04;. Found: C, 68.64; H, 6.08.

Example 529,9,9-trifluoro-8-oxo-N-(4'-(trifluoromethoxy)(1,1'-biphenyl)-3-yl)nonanamide

The desired product was prepared by substituting 4'-(trifluoromethoxy)(1,1'-biphenyl)-3-amine for aniline in Example 1. MS (DCI) m/e 462 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.98 (s, 1H), 7.94 (s, 1H), 7.63 (d, 2H), 7.57 (d, 1H), 7.47 (d, 2H), 7.38 (d, 1H), 7.34 (d, 1H), 2.86 (t, 2H), 2.33 (t, 2H), 1.68-1.50 (m, 4H), 1.40-1.25 (m, 4H); Anal. Calcd for C₂₂H₂₁NO₃F₆: C, 57.26; H, 4.58; N, 3.03. Found: C, 57.03; H, 4.65; N, 2.92.

10

Example 539,9,9-trifluoro-8-oxo-N-(3-(3-pyridinyl)phenyl)nonanamide

The desired product was prepared by substituting 3-(3-pyridinyl)aniline for aniline in Example 1. MS (DCI) m/e 379 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 10.0 (s, 1H), 8.82 (s, 1H), 8.58 (d, 1H), 8.00 (d, 1H), 7.95 (s, 1H), 7.64 (d, 1H), 7.53-7.31 (m, 3H), 2.86 (t, 2H), 2.33 (t, 2H), 1.69-1.52 (m, 4H), 1.41-1.20 (m, 4H); Anal. Calcd for C₂₀H₂₁N₂O₂F₃·0.5H₂O: C, 62.01; H, 5.72; N, 7.23. Found: C, 61.98; H, 5.73; N, 7.05.

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Example 549,9,9-trifluoro-N-(4'-(methylsulfanyl)(1,1'-biphenyl)-3-yl)-8-oxononanamide

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Example 54AN-(3-bromophenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 3-bromoaniline for aniline in Example 1. MS (ESI(+)) m/e 381 (M+H)⁺.

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Example 54B9,9,9-trifluoro-N-(4'-(methylsulfanyl)(1,1'-biphenyl)-3-yl)-8-oxononanamide

A mixture of Example 54A (308 mg, 0.81 mmol), 4-(methylsulfanyl)phenyl-boronic acid (150 mg, 0.89 mmol), Pd(OAc)₂ (9.1 mg, 0.04 mmol), tri-o-tolylphosphine (24.4 mg, 0.08 mmol), and 2M Na₂CO₃ (2 mL, 2mmol) in DME (5 mL) was heated to 80 °C for 3 hours, treated with additional Pd (OAc)₂ (9 mg), tri-o-tolylphosphine (24 mg), and 3-(methylsulfanyl)phenyl-boronic acid (75 mg), heated for 3 hours, cooled to room temperature, and partitioned between diethyl ether and water. The aqueous phase was extracted with diethyl ether and the combined extracts were washed with brine, dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 7:3 hexanes/ethyl acetate to provide 164 mg (48%) of the desired product. MS (ESI(-)) m/e 422 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.94 (s,

1H), 7.90 (s, 1H), 7.55 (d, 3H), 7.42-7.22 (m, 4H), 2.86 (t, 2H), 2.49 (s, 3H), 2.33 (t, 2H), 1.68-1.51 (m, 4H), 1.36-1.27 (m, 4H); Anal. Calcd for $C_{22}H_{24}NO_2F_3S$: C, 62.39; H, 5.71; N, 3.31. Found: C, 63.18; H, 5.60; N, 2.75.

5

Example 55

N-(3'-amino(1,1'-biphenyl)-3-yl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 3-aminophenylboronic acid for 4-(methylsulfanyl)phenylboronic acid in Example 54. MS (ESI(-)) m/e 391 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.92 (s, 1H), 7.87 (s, 1H), 7.50 (d, 1H), 7.20 (d, 1H), 7.09 (t, 1H), 6.78 (dd, 1H), 6.72 (d, 1H), 6.56 (d, 1H), 5.17 (s, 2H), 2.87 (t, 2H), 2.34 (t, 2H), 1.67-1.52 (m, 4H), 1.38-1.27 (m, 4H); Anal. Calcd for $C_{21}H_{23}N_2O_2F_3$: C, 64.28; H, 5.91; N, 7.14. Found: C, 63.95; H, 5.99; N, 6.91.

Example 56

9,9,9-trifluoro-N-(4'-(methylsulfonyl)(1,1'-biphenyl)-3-yl)-8-oxononanamide

A suspension of Example 54B (85 mg, 0.2 mmol) in a 2:1 mixture of methanol/water (10 mL) at room temperature was treated with NaHCO₃ (42 mg, 0.5 mmol) and oxone (10 mL), stirred for 18 hours, and partitioned between ethyl acetate and water. The aqueous phase was extracted with ethyl acetate and the combined extracts were washed with brine, dried (Na₂SO₄), filtered, and concentrated. The concentrate was recrystallized from ethyl acetate/hexane to provide 56 mg (62%) of the desired product. MS (ESI(-)) m/e 454 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 10.03 (s, 1H), 8.03 (d, 3H), 7.86 (d, 2H), 7.64 (d, 1H), 7.48-7.38 (m, 2H), 3.27 (s, 3H), 2.87 (t, 2H), 2.33 (t, 2H), 1.67-1.52 (m, 4H), 1.38-1.27 (m, 4H); Anal. Calcd for $C_{22}H_{24}NO_4F_3S \cdot 0.5H_2O$: C, 56.89; H, 5.42; N, 3.02. Found: C, 56.80; H, 5.47; N, 2.82.

Example 57

N-(4'-cyano(1,1'-biphenyl)-3-yl)-8,8,8-trifluoro-7-oxooctanamide

The desired product was prepared by substituting methyl 8,8,8-trifluoro-7-oxooctanoate and 3'-amino(1,1'-biphenyl)-4-carbonitrile for Example 1A and aniline, respectively, in Example 1. MS (ESI(-)) m/e 387 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 10.04 (s, 1H), 8.03 (s, 1H), 7.93 (d, 2H), 7.80 (d, 2H), 7.63 (d, 1H), 7.48-7.35 (m, 2H), 2.87 (t, 2H), 2.33 (t, 2H), 1.69-1.52 (m, 4H), 1.43-1.23 (m, 2H); Anal. Calcd for $C_{21}H_{19}N_2O_2F_3 \cdot 0.25H_2O$: C, 64.20; H, 5.0; N, 7.13. Found: C, 64.26; H, 5.08; N, 7.13.

35

Example 58

N-(3-cyanophenyl)-8,8,8-trifluoro-7-oxooctanamide

The desired product was prepared by substituting methyl 8,8,8-trifluoro-7-oxooctanoate and 3-cyanoaniline for Example 1A and aniline, respectively, in Example 1. MS (ESI(-)) m/e 311 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 10.22 (s, 1H), 8.10 (s, 1H), 7.78 (dd, 1H), 7.56-7.43 (m, 2H), 2.88 (t, 2H), 2.33 (t, 2H), 1.68-1.53 (m, 2H), 1.51-1.20 (m, 4H); Anal. Calcd for C₁₅H₁₅N₂O₂F₃·0.65H₂O: C, 55.61; H, 5.07; N, 8.65. Found: C, 55.67; H, 5.0; N, 8.51.

Example 59

8,8,8-trifluoro-7-oxo-N-(4-phenoxyphenyl)octanamide

The desired product was prepared by substituting methyl 8,8,8-trifluoro-7-oxooctanoate and 4-phenoxyaniline for Example 1A and aniline, respectively, in Example 1. MS (ESI(-)) m/e 378 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.88 (s, 1H), 7.60 (d, 2H), 7.36 (t, 2H), 7.08 (t, 1H), 6.97 (t, 4H), 2.88 (t, 2H), 2.28 (t, 2H), 1.67-1.52 (m, 4H), 1.41-1.28 (m, 2H); Anal. Calcd for C₂₀H₂₀NO₃F₃: C, 63.32; H, 5.31; N, 3.69. Found: C, 63.14; H, 5.22; N, 3.53.

Example 60

N-(4-aminobenzyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 4-(aminomethyl)aniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 329 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 8.06 (t, 1H), 7.22 (dt, 1H), 7.12 (d, 1H), 7.00 (d, 1H), 6.88 (t, 1H), 4.20 (d, 2H), 2.14 (t, 2H), 1.60 (m, 2H), 1.52 (m, 2H), 1.42 (m, 2H), 1.22 (m, 4H).

Example 61

9,9,9-trifluoro-N-(3-methylphenyl)-8-oxononanamide

The desired product was prepared by substituting 3-methylaniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 314 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.80 (s, 1H), 7.44 (s, 1H), 7.36 (d, 1H), 7.15 (t, 1H), 6.85 (d, 1H), 2.28 (t, 2H), 2.26 (s, 3H), 1.6 (m, 4H), 1.44 (m, 2H), 1.28 (m, 4H).

30

Example 62

9,9,9-trifluoro-N-(4-methylphenyl)-8-oxononanamide

The desired product is prepared by substituting 4-methylaniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 314 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.74 (s, 1H), 7.42 (d, 2H), 7.04 (d, 2H), 2.28 (t, 2H), 2.26 (s, 3H), 1.62 (m, 4H), 1.22 (m, 6H).

Example 63

N-(4-aminophenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 1,4-benzenediamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 315 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.80 (s, 1H), 7.45 (d, 2H), 7.0 (d, 2H), 6.60 (br s, 2H), 2.24 (t, 2H), 1.46 (m, 4H), 1.40 (m, 2H), 1.22 (m, 4H).

Example 649,9,9-trifluoro-N-(4-fluorobenzyl)-8-oxononanamide

The desired product was prepared by substituting (4-fluorophenyl)methanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 332 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.02 (t, 0.54H), 8.24 (t, 0.46H), 8.00 (m, 1H), 7.35 (m, 1H), 7.30 (m, 1H), 7.24 (m, 1H), 4.42 (d, 1.08H), 4.21 (d, 0.96H), 2.10 (dt, 2H), 1.60 (m, 2H), 1.52 (m, 2H), 1.42 (m, 2H), 1.24 (m, 4H).

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Example 659,9,9-trifluoro-N-(3-methoxybenzyl)-8-oxononanamide

The desired product was prepared by substituting (3-methoxyphenyl)methanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 344 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.00 (t, 1H), 7.22 (t, 1H), 7.05 (dd, 1H), 6.90 (d, 1H), 6.80 (d, 1H), 4.40 (d, 2H), 3.82 (s, 3H), 2.24 (t, 2H), 1.60 (m, 2H), 1.52 (m, 2H), 1.42 (m, 2H), 1.24 (m, 4H).

Example 669,9,9-trifluoro-N-(4-methoxybenzyl)-8-oxononanamide

The desired product was prepared by substituting (4-methoxyphenyl)methanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 344 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 8.18 (t, 1H), 7.40 (d, 1H), 7.10 (d, 1H), 7.00 (d, 1H), 6.80 (d, 1H), 4.20 (d, 2H), 3.70 (s, 3H), 2.10 (t, 2H), 1.60 (m, 2H), 1.50 (m, 2H), 1.40 (m, 2H), 1.24 (m, 4H).

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Example 679,9,9-trifluoro-N-(3-fluorobenzyl)-8-oxononanamide

The desired product was prepared by substituting (3-fluorophenyl)methanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 332 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.20 (t, 0.5H), 8.30 (t, 0.5H), 7.78 (dd, 0.5H), 7.65 (dd, 0.5H), 7.54 (m, 1H), 7.36 (m, 1H), 7.15 (m, 0.5H), 7.05 (m, 0.5H), 4.45 (d, 1H), 4.22 (d, 1H), 2.30 (t, 1H), 2.14 (t, 1H), 1.60 (m, 2H), 1.52 (m, 2H), 1.42 (m, 2H), 1.24 (m, 4H).

Example 68

9,9,9-trifluoro-N-(3-chlorobenzyl)-8-oxononanamide

The desired product was prepared by substituting (3-chlorophenyl)methanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 348 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 8.30 (t, 1H), 7.34 (m, 1H), 7.28 (m, 2H), 7.20 (d, 1H), 4.20 (d, 2H), 2.16 (t, 2H), 1.60 (m, 2H), 1.50 (m, 2H), 1.40 (m, 2H), 1.22 (m, 4H).

Example 69N-(4-bromobenzyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting (4-bromophenyl)methanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 392 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 8.30 (t, 1H), 7.50 (d, 2H), 7.20 (d, 2H), 4.20 (d, 2H), 2.12 (m, 2H), 1.60 (m, 2H), 1.50 (m, 2H), 1.40 (m, 2H), 1.22 (m, 4H).

Example 70N-(3-(dimethylamino)phenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting N,N-dimethyl-1,3-benzenediamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 343 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.60 (d, 1H), 7.05 (m, 2H), 6.90 (m, 1H), 6.41 (d, 1H), 2.50 (s, 6H), 2.20 (m, 2H), 1.60 (m, 2H), 1.50 (m, 2H), 1.40 (m, 2H), 1.22 (m, 4H).

Example 719,9,9-trifluoro-8-oxo-N-(3-(trifluoromethoxy)benzyl)nonanamide

The desired product was prepared by substituting (3-(trifluoromethoxy)phenyl)methanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 398 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 8.38 (t, 1H), 7.40 (t, 1H), 7.25 (d, 1H), 7.20 (m, 2H), 4.25 (d, 2H), 2.10 (t, 2H), 1.60 (m, 1H), 1.48 (m, 4H), 1.40 (m, 1H), 1.22 (m, 4H).

Example 729,9,9-trifluoro-8-oxo-N-(3-(trifluoromethyl)benzyl)nonanamide

The desired product was prepared by substituting (3-(trifluoromethyl)phenyl)methanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 382 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 8.40 (t, 1H), 7.60 (m, 4H), 4.36 (d, 2H), 2.14 (m, 2H), 1.60 (m, 4H), 1.40 (m, 1H), 1.22 (m, 4H).

Example 739,9,9-trifluoro-8-oxo-N-(3-(trifluoromethoxy)phenyl)nonanamide

The desired product was prepared by substituting 3-(trifluoromethoxy)aniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 384 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 10.10 (s, 1H), 7.52 (d, 1H), 7.40 (m, 2H), 7.0 (d, 1H), 2.30 (m, 2H), 1.60 (m, 5H), 1.40 (m, 1H), 1.20 (m, 4H).

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Example 74

N-(3,5-dimethoxybenzyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting (3,5-dimethoxyphenyl)-methanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 374 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 8.20 (s, 1H), 6.56 (s, 1H), 6.40 (s, 2H), 4.20 (d, 2H), 3.60 (s, 6H), 2.12 (t, 2H), 1.60 (m, 1H), 1.54 (m, 4H), 1.40 (m, 1H), 1.22 (m, 4H).

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Example 75

N-(2,4-dimethylphenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 2,4-dimethylaniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 328 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.10 (s, 1H), 7.20 (d, 1H), 7.00 (s, 1H), 6.95 (d, 1H), 2.24 (t, 2H), 2.20 (s, 3H), 2.14 (s, 3H), 1.60-1.40 (m, 6H), 1.28-1.20 (m, 4H).

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Example 76

N-(3,4-dimethylphenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 3,4-dimethylaniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 328 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.60 (s, 1H), 7.40 (s, 1H), 7.34 (d, 1H), 7.0 (d, 1H), 2.24 (t, 2H), 2.18 (s, 3H), 2.12 (s, 3H), 1.60 (m, 5H), 1.40 (m, 1H), 1.28 (m, 4H).

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Example 77

N-(3,5-dimethylphenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 3,5-dimethylaniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 328 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.60 (s, 1H), 7.20 (s, 2H), 6.64 (s, 1H), 2.24 (t, 2H), 2.20 (s, 6H), 1.60 (m, 5H), 1.40 (m, 1H), 1.26 (m, 4H).

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Example 78

N-(2,4-dimethoxyphenyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 2,4-dimethoxyaniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 360 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆)

δ 8.82 (s, 1H), 7.60 (d, 1H), 6.60 (s, 1H), 6.42 (d, 1H), 3.66 (s, 3H), 3.62 (s, 3H), 2.30 (t, 2H), 1.60 (m, 5H), 1.40 (m, 1H), 1.26 (m, 4H).

Example 79

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N-(2,5-dimethoxybenzyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting (2,5-dimethoxyphenyl)methanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 374 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 8.12 (t, 1H), 6.90 (d, 1H), 6.68 (d, 1H), 6.62 (s, 1H), 4.2 (d, 2H), 3.72 (s, 3H), 3.64 (s, 3H), 2.10 (t, 2H), 1.60-1.40 (m, 6H), 1.20 (m, 4H).

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Example 80

N-(3,5-dimethoxyphenyl)-9,9,9-trifluoro-8-oxononanamide

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The desired product was prepared by substituting 3,5-dimethoxyaniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 360 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.80 (s, 1H), 6.80 (s, 2H), 6.20 (s, 1H), 3.70 (s, 6H), 2.24 (t, 2H), 1.60 (m, 4H), 1.40 (m, 2H), 1.22 (m, 4H).

Example 81

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N-(1,3-benzodioxol-5-ylmethyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 1,3-benzodioxol-5-ylmethanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 358 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 8.80 (t, 1H), 6.86 (d, 1H), 6.82 (s, 1H), 6.76 (d, 1H), 6.0 (s, 2H), 4.20 (d, 2H), 2.10 (t, 2H), 1.80-1.40 (m, 6H), 1.20 (m, 4H).

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Example 82

9,9,9-trifluoro-8-oxo-N-(3,4,5-trimethoxyphenyl)nonanamide

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The desired product was prepared by substituting 3,4,5-trimethoxyaniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 390 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 9.70 (s, 1H), 7.00 (s, 2H), 3.80 (s, 6H), 3.60 (s, 3H), 2.22 (t, 2H), 1.60 (m, 4H), 1.20 (m, 2H), 1.20 (m, 4H).

Example 83

N-(3,4-dichlorobenzyl)-9,9,9-trifluoro-8-oxononanamide

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The desired product was prepared by substituting (3,4-dichlorophenyl)-methanamine for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 382 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 8.40 (t, 1H), 7.60 (d, 1H), 7.42 (s, 1H), 7.20 (d, 1H), 4.22 (d, 2H), 2.12 (t, 2H), 1.62-1.40 (m, 6H), 1.20 (m, 4H).

Example 848-((1,1'-biphenyl)-4-yloxy)-1,1,1-trifluoro-3-octyn-2-one

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Example 84A4-(5-hexynyloxy)-1,1'-biphenyl

A solution of (1,1'-biphenyl)-4-ol (1.70 g, 10.0 mmol), 5-hexyn-1-ol (0.98 g, 10.0 mmol), and triphenylphosphine (3.41 g, 13.0 mmol) in THF (20 mL) at 0 °C was treated dropwise with diethylazodicarboxylate (2.27 g, 13.0 mmol), warmed to room temperature, 10 stirred for 18 hours, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 50:1 hexanes/ethyl acetate to provide 1.68 g (67%) of the desired product.

Example 84B8-((1,1'-biphenyl)-4-yloxy)-1,1,1-trifluoro-3-octyn-2-one

A solution of Example 84A (1.51 g, 6.0 mmol) in THF (35 mL) at -78 °C was treated dropwise with 2.5 M n-butyllithium in hexanes (2.6 mL, 6.5 mmol) and ethyl trifluoroacetate (0.98 g, 6.9 mmol), stirred for 10 minutes, treated with boron trifluoride diethyl etherate (1.50 g, 10 mmol), stirred for 4 hours, warmed to room temperature, and stirred for 18 hours. The 20 mixture was quenched with saturated NH₄Cl (20 mL), and extracted with diethyl ether. The combined extracts were washed with brine, dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 95:5 to 92:8 hexanes/ethyl acetate to provide 959 mg (46%) of the desired product. MS (ESI(-)) m/e 345 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 7.62-7.57 (m, 4H), 7.45-7.40 (m, 2H), 7.33-7.27 (m, 1H), 7.02 (d, 2H), 4.06 (t, 2H), 2.78 (t, 2H), 1.91-1.71 (m, 4H); Anal. Calcd for 25 C₂₀H₁₇F₃O₂: C, 69.36; H, 4.95. Found: C, 69.45; H, 5.08.

Example 85N-(4-(dimethylamino)benzyl)-9,9,9-trifluoro-8-oxononanamide

The desired product was prepared by substituting 4-(aminomethyl)-N,N-dimethylaniline for 4-aminopyridine in Example 6. MS (ESI(-)) m/e 357 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 8.06 (t, 1H), 7.12 (d, 2H), 6.88 (d, 2H), 4.20 (d, 2H), 2.80 (s, 6H), 2.14 (t, 2H), 1.60 (m, 2H), 1.52 (m, 2H), 1.42 (m, 2H), 1.22 (m, 4H).

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Example 86(3E)-8-((1,1'-biphenyl)-4-yloxy)-1,1,1-trifluoro-3-octen-2-one

Example 86A(3E)-8-((1,1'-biphenyl)-4-yloxy)-1,1,1-trifluoro-3-octen-2-ol

A suspension of LAH (201 mg, 5.3 mmol) in THF (10 mL) at 0 °C was treated dropwise with a solution of Example 84B (822 mg, 2.40 mmol) in THF (2.5 mL), stirred for 5 30 minutes, warmed to room temperature, heated to reflux for 8 hours, cooled to 0 °C, and treated sequentially with water (0.1 mL), 1M NaOH (0.1 mL), and water (0.5 mL). The suspension was filtered and the filtrate was washed sequentially with saturated NH₄Cl, water, and brine, dried (MgSO₄), filtered, and concentrated to provide 660 mg (78%) of the desired product. MS (ESI(+)) m/e 368 (M+NH₄)⁺.

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Example 86B(3E)-8-((1,1'-biphenyl)-4-yloxy)-1,1,1-trifluoro-3-octen-2-one

A solution of Example 86A (201 mg, 0.57 mmol) in dichloromethane (3 mL) at room temperature was added dropwise to a suspension of Dess-Martin reagent (894 mg, 2.10 15 mmol) in dichloromethane (20 mL), stirred for 3 hours, diluted with 1M NaOH (20 mL) and diethyl ether (20 mL), and stirred for 30 minutes. The aqueous phase was extracted with diethyl ether and the combined organic phases were washed with brine, dried (MgSO₄), filtered, and concentrated to provide 180 mg (90%) of the desired product. MS (ESI(-)) m/e 347 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 7.62-7.53 (m, 4H), 7.45-7.27 (m, 4H), 7.01 20 (d, 2H), 6.67 (dd, 1H), 4.03 (t, 2H), 2.49-2.43 (m, 2H), 1.82-1.61 (m, 4H); Anal. Calcd for C₂₀H₁₉F₃O₂: C, 68.96; H, 5.50. Found: C, 69.11; H, 5.42.

Example 87(8E)-9-(1,1'-biphenyl)-4-yl-1,1,1-trifluoro-8-nonen-2-one

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Example 87Aethyl (7E)-8-(1,1'-biphenyl)-4-yl-7-octenoate

A solution of (7-ethoxy-7-oxoheptyl)(triphenyl)phosphonium bromide (599 mg, 1.2 30 mmol) in THF (5mL) at 0 °C was treated with potassium tert-butoxide (115 mg, 1.2 mmol) and 4-phenylbenzaldehyde (182 mg, 1.0 mmol), stirred for 1.5 hours, warmed to room temperature, and treated with ethyl acetate and water. The aqueous phase was extracted with ethyl acetate, and the combined organic phases were washed with brine, dried (MgSO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 5% ethyl acetate/hexanes to provide 244mg (76%) of the desired product. MS 35 (ESI(+)) m/e 323 (M+H)⁺.

Example 87B

(8E)-9-(1,1'-biphenyl)-4-yl-1,1,1-trifluoro-8-nonen-2-one

The desired product was prepared by substituting Example 87A for Example 2A in Examples 2B and 2C. MS (ESI(-)) m/e 345 (M-H)⁻; ¹H NMR (300 MHz, DMSO-d₆) δ 7.80-7.30 (m, 9H), 6.45 (br d, 1H), 5.69 (dt, 1H), 2.87 (t, 2H), 2.35 (dq, 2H), 1.60-1.50 (m, 2H), 1.50-1.40 (m, 2H), 1.40-1.30 (m, 2H); Anal. Calcd for C₂₁H₂₁F₃O·0.7H₂O: C, 70.61; H, 6.26. Found: C, 70.53; H, 5.76.

Example 881-(2-(4-((1,1'-biphenyl)-4-yloxy)butyl)cyclopropyl)-2,2,2-trifluoroethanone

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Example 88A1-(2-(4-((1,1'-biphenyl)-4-yloxy)butyl)cyclopropyl)-2,2,2-trifluoroethanol

Samarium (362 mg, 2.4 mmol) was dried under vacuum with heating, purged with nitrogen, treated with THF (3 mL) and a solution of Example 87A (200 mg, 0.57 mmol) in THF (4 mL), cooled to 0 °C, and treated with CH₂I₂ (0.18 mL, 2.2 mmol). The reaction was warmed to room temperature, stirred for 24 hours, and partitioned between saturated K₂CO₃ and diethyl ether. The organic phase was washed with brine, dried (MgSO₄), filtered, and concentrated. The concentrate was purified by HPLC with 20% ethyl acetate/hexanes to provide 130 mg (63%) of the desired product. MS (APCI(+)) m/e 382 (M+NH₄)⁺.

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Example 88B1-(2-(4-((1,1'-biphenyl)-4-yloxy)butyl)cyclopropyl)-2,2,2-trifluoroethanone

The desired product was prepared by substituting Example 88A for Example 86A in Example 86B. MS (ESI(+)) m/e 380 (M+NH₄)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 7.62-7.56 (m, 4H), 7.45-7.39 (m, 2H), 7.33-7.27 (m, 1H), 7.00 (d, 2H), 4.01 (t, 2H), 2.36-2.31 (m, 1H), 1.77-1.71 (m, 2H), 1.70-1.62 (m, 1H), 1.59-1.40 (m, 5H), 1.33-1.27 (m, 1H); Anal. Calcd for C₂₁H₂₁F₃O₂·0.3H₂O: C, 68.58; H, 5.92. Found: C, 68.37; H, 5.62.

Example 89

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9H-fluoren-9-ylmethyl 7,7,7-trifluoro-6-oxoheptylcarbamate

A suspension of Fmoc-ε-aminocaproic acid (4.97g, 14.1 mmol) in dichloromethane (25 mL) at room temperature was treated with oxalyl chloride (2.2 mL, 25.2 mmol), stirred for 1 hour, and concentrated. The concentrate was dissolved in dichloromethane (100 mL), treated with trifluoroacetic anhydride (8.92g, 42.4 mmol), cooled to -50 °C, and treated dropwise with pyridine (5.7 mL, 70 mmol) over 5 minutes. The mixture was warmed to -25 °C, stirred for 1 hour, warmed to 10 °C over 1 hour, cooled to -30 °C, quenched with water (10 mL), and partitioned between dichloromethane and water. The organic phase was

washed with water, dried (Na_2SO_4), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 30% ethyl acetate/hexanes to 50% ethyl acetate/hexanes to 80% ethyl acetate/hexanes to provide 900 mg (16%) of the desired product. mp: 83-88 °C; MS (ESI(+)) m/e 406 ($\text{M}+\text{H}$)⁺; ^1H NMR (300 MHz, CDCl_3) δ 1.39-5 1.27 (m, 2H), 1.56-1.43 (m, 2H), 1.63-1.58 (m, 2H), 2.68 (t, 2H), 3.09-2.92 (m, ~0.5H), 3.17 (q, 2H), 4.20 (t, 1H), 4.40 (d, 2H), 4.54-4.43 (m, ~0.5H), 4.89-4.76 (m, 1H), 7.28 (td, 2H), 7.37 (t, 2H), 7.57 (d, 2H), 7.73 (d, 2H); Anal. Calcd for $\text{C}_{22}\text{H}_{22}\text{F}_3\text{NO}_3$: C, 65.18; H, 5.47; N, 3.45; F, 14.06. Found: C, 64.94; H, 5.53; N, 3.46; F, 13.75.

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Example 904-((1,1'-biphenyl)-4-yloxy)-N-(3,3,3-trifluoro-2-oxopropyl)butanamideExample 90Amethyl 4-((1,1'-biphenyl)-4-yloxy)butanoate

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A mixture of (1,1'-biphenyl)-4-ol (2.0 g, 11.75 mmol), methyl 4-bromobutyrate (1.55 mL, 12.0 mmol) and Cs_2CO_3 (4.21 g, 12.9 mmol) in DMF (40 mL) at room temperature was stirred for 18 hours, diluted with water (500 mL), and filtered to provide 2.98 g (94%) of the desired product. MS (ESI(+)) m/e 271 ($\text{M}+\text{H}$)⁺.

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Example 90Blithium 4-((1,1'-biphenyl)-4-yloxy)butanoate

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A mixture of Example 90A (2.0 g, 8.0 mmol), 2M LiOH in water (24 mL, 48 mmol) and THF (18 mL) at room temperature was stirred for 18 hours, partially concentrated, and filtered. The solid was washed with water and dried in a vacuum oven to provide 1.94 g (93%) of the desired product. MS (ESI(-)) m/e 255 ($\text{M}-\text{Li}$)⁻.

Example 90C4-((1,1'-biphenyl)-4-yloxy)-N-(3,3,3-trifluoro-2-hydroxypropyl)butanamide

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A mixture of Example 90B (206 mg, 0.8 mmol), 2-hydroxy-2-

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(trifluoromethyl)ethylamine (102 mg, 0.79 mmol, prepared as described in J.Org.Chem. 1995, 60, 41), EDCI (162 mg, 0.85 mmol), HOBr (115 mg, 0.85 mmol) and NMM (0.17 mL, 1.5 mmol) in DMF (2 mL) at room temperature was stirred for 18 hours, and partitioned between ethyl acetate and water. The organic phase was washed with brine, dried (Na_2SO_4), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 20% ethyl acetate/hexanes to 30% ethyl acetate/hexanes to 40% ethyl acetate/hexanes to provide 0.16 g (55%) of the desired product. MS (ESI(-)) m/e 366 ($\text{M}-\text{H}$)⁻.

Example 90D4-((1,1'-biphenyl)-4-yloxy)-N-(3,3,3-trifluoro-2-oxopropyl)butanamide

The desired product was prepared by substituting Example 90C for Example 86A in Example 86B. MS (ESI(+)) m/e 366 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.08-8.04 (m, 1H), 7.62-7.57 (m, 4H), 7.45-7.40 (m, 2H), 7.32-7.27 (m, 1H), 7.00 (d, 2H), 4.01 (t, 2H), 3.40 (d, 2H), 2.34 (t, 2H), 1.99-1.93 (m, 2H); Anal. Calcd for C₁₉H₁₈F₃NO₃·0.8H₂O: C, 60.09; H, 5.20; N, 3.69. Found: C, 59.74; H, 5.35; N, 3.51.

Example 91methyl 8-((1,1'-biphenyl)-4-yloxy)-2-oxooctanoateExample 91A6-((4'-phenylphenoxy))-hexan-1-ol

A solution of (1,1'-biphenyl)-4-ol (2.3 g, 13.5 mmol) in DMF (15 mL) at room temperature was treated with Cs₂CO₃ (8.8 g, 27 mmol), stirred for 20 minutes, treated with a solution of 6-(t-butyldimethylsilyloxy)hexyl bromide (4.78 g, 16.2 mmol) in DMF (5 mL), stirred for 48 hours, and partitioned between water and diethyl ether. The aqueous phase was extracted with diethyl ether, and the combined organic phases were washed with brine, dried (MgSO₄), filtered, and concentrated. The concentrate was dissolved in THF (60 mL), treated with 1M TBAF in THF (27 mL, 27 mmol), stirred for 18 hours, poured into water, and extracted with ethyl acetate. The combined extracts were washed with brine, dried (MgSO₄), filtered, and concentrated. The concentrate was triturated with hexanes and filtered to provide the desired product.

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Example 91B6-((1,1'-biphenyl)-4-yloxy)hexanal

A solution of oxalyl chloride (1.24 mL, 14.22 mmol) in dichloromethane (10 mL) at -60 °C was treated dropwise with a solution of DMSO (1.85 mL, 26.1 mmol) in dichloromethane (2 mL), stirred for 10 minutes, treated with a solution of Example 91A (3.2 g, 11.85 mmol) in dichloromethane (10 mL), stirred for 15 minutes, treated with triethylamine (8.19 mL, 59.2 mmol), stirred for 5 minutes, warmed to room temperature, and partitioned between water and dichloromethane. The organic extract was washed with brine, dried (MgSO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 10% ethyl acetate/hexanes to provide the desired product.

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Example 91Cmethyl 8-((1,1'-biphenyl)-4-yloxy)-2-oxooctanoate

A suspension of lithium chloride (24 mg, 0.58 mmol) in acetonitrile (2 mL) at room temperature was treated with a solution of methyl (dimethoxyphosphoryl)-(tetrahydro-2H-pyran-2-yloxy)acetate (150 mg, 0.53 mmol, prepared according to the procedure described in *Tet. Lett.* **1981**, 22, 663-666) in acetonitrile (1.5 mL), treated with DBU (0.07 mL, 0.47 mmol), stirred for 10 minutes, cooled to 0 °C, treated with a solution of Example 91B (118 mg, 0.44 mmol) in acetonitrile (2 mL), stirred for 1.5 hours, and warmed to room temperature. The reaction was partitioned between water and diethyl ether and the organic extract was dried (MgSO_4), filtered, and concentrated. The concentrate was dissolved in methanol (10 mL), treated with $\text{pTsOH}\cdot\text{H}_2\text{O}$ (15 mg), stirred for 45 minutes, and concentrated. The concentrate was dissolved in dichloromethane, washed with saturated NaHCO_3 , dried (Na_2SO_4), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 10% ethyl acetate/hexanes to provide 63 mg (42%) of the desired product. MS (ESI(+)) m/e 358 ($\text{M}+\text{NH}_4$)⁺; ¹H NMR (300 MHz, DMSO-d_6) δ 7.62-7.56 (m, 4H), 7.45-7.40 (m, 2H), 7.33-7.28 (m, 1H), 7.01 (d, 2H), 4.00 (t, 2H), 3.32 (s, 3H), 2.83 (t, 2H), 1.77-1.69 (m, 2H), 1.58-1.49 (m, 2H), 1.48-1.32 (m, 4H); Anal. Calcd. for $\text{C}_{21}\text{H}_{24}\text{O}_4$: C, 74.09; H, 7.11. Found: C, 74.07; H, 6.93.

Example 92

7-((1,1'-biphenyl)-3-yloxy)-1-(1,3-oxazol-2-yl)-1-heptanone

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Example 92A

ethyl 7-((1,1'-biphenyl)-3-yloxy)heptanoate

The desired product was prepared by substituting (1,1'-biphenyl)-3-ol for (1,1'-biphenyl)-4-ol in Example 2A.

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Example 92B

7-((1,1'-biphenyl)-3-yloxy)heptanoic acid

The desired product was prepared by substituting Example 92A for Example 1A in Example 1B.

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Example 92C

7-((1,1'-biphenyl)-3-yloxy)heptanoyl chloride

The desired product was prepared by substituting Example 92B for Example 102B in Example 102C.

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Example 92D

7-((1,1'-biphenyl)-3-yloxy)-1-(1,3-oxazol-2-yl)-1-heptanone

A solution of oxazole (39 mg, 0.56 mmol) in THF (6 mL) at -78 °C was treated dropwise with 2.5M n-butyllithium in hexanes (0.34 mL, 0.85 mmol), stirred for 20 minutes, treated with 0.5M ZnCl₂ in THF (2.26 mL, 1.13 mmol), warmed to 0 °C, and stirred for 45 minutes. The mixture was treated with CuI (107 mg, 0.56 mmol), stirred for 10 minutes, 5 treated dropwise with a solution of Example 92C (1.13 mmol) in THF (4 mL), and stirred for 1 hour. The mixture was treated with ethyl acetate (30 mL), washed sequentially with 15% NH₄OH (20 mL), water (20 mL), and saturated NH₄Cl (10 mL), dried (MgSO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 20 % ethyl acetate/hexanes to provide the desired product. MS (ESI(+)) m/e 350 10 (M+H)⁺; ¹H NMR (CDCl₃) δ 7.80 (s, 1H), 7.60 (m, 1H), 7.58 (m, 1H), 7.42 (m, 2H), 7.36 (m, 1H), 7.34 (m, 1H), 7.32 (s, 1H), 7.16 (m, 1H), 7.10 (t, 1H), 6.88 (dd, 1H), 4.00 (t, 2H), 3.10 (t, 2H), 1.85-1.75 (m, 4H), 1.55-1.45 (m, 4H).

Example 93

8-((1,1'-biphenyl)-4-yloxy)-2-oxooctanoic acid

The desired product was prepared by substituting Example 91C for Example 1A in Example 1B. MS (ESI(+)) m/e 344 (M+NH₄)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 7.62-7.56 (m, 4H), 7.45-7.40 (m, 2H), 7.32-7.27 (m, 1H), 7.03-7.00 (m, 2H), 4.00 (t, 2H), 2.66 (t, 2H), 1.77-1.67 (m, 2H), 1.56-1.24 (m, 6H); Anal. Calcd. for C₂₀H₂₂O₄·H₂O: C, 69.75; H, 7.02. 20 Found: C, 69.76; H, 6.70.

Example 94

ethyl 7-((1,1'-biphenyl)-4-yloxy)-2-oxoheptanoate

Example 94A

ethyl 7-((1,1'-biphenyl)-4-yloxy)-2-hydroxyheptanoate

A solution of 0.5M KHMDS in THF (9.2 mL, 4.6 mmol) in THF (100 mL) at -78 °C was treated with a solution of Example 2A (1.0 g, 3.0 mmol) in THF (50 mL), stirred for 15 minutes, treated with 2-benzenesulfonyl-3-phenyl-oxaziridine (1.2 g, 4.6 mmol, 30 prepared according to the procedure described in J. Org. Chem. 1982, 47, 1774-1775) stirred for 30 minutes, quenched with saturated NH₄Cl, and extracted with ethyl acetate. The combined extracts were dried (MgSO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 4:1 ethyl acetate/dichloromethane to provide 0.51 g (50%) of the desired product.

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Example 94B

ethyl 7-((1,1'-biphenyl)-4-yloxy)-2-oxoheptanoate

A solution of Example 94A (155 mg, 0.45 mmol) in dichloromethane (3 mL) at 0 °C was treated sequentially with 4A molecular sieves and PDC (256 mg, 0.68 mmol), warmed to room temperature, stirred for 72 hours, diluted with ethyl acetate, filtered through diatomaceous earth (Celite®), and concentrated. The concentrate was purified by flash column chromatography on silica gel with 8:1 ethyl acetate/hexanes to provide the desired product. MS (DCI/NH₃) m/e 358 (M+NH₄)⁺; ¹H NMR (300 MHz, CDCl₃) δ 7.56-7.49 (m, 4H), 7.44-7.38 (m, 2H), 7.32-7.27 (m, 1H), 6.98-6.93 (m, 2H), 4.32 (q, 2H), 4.00 (t, 2H), 2.89 (t, 2H), 1.88-1.69 (m, 4H), 1.60-1.55 (m, 2H), 1.37 (t, 3H); Anal. Calcd. for C₂₁H₂₄O₄: C, 74.09; H, 7.11. Found: C, 74.10; H, 7.03.

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Example 95ethyl 7-((3-bromo(1,1'-biphenyl)-4-yl)oxy)-2-oxoheptanoateExample 95A4-((5-bromopentyl)oxy)-1,1'-biphenyl

A solution of 1,5-dibromopentane (6.4 mL, 50 mmol) and (1,1'-biphenyl)-4-ol (2.67 g, 15.7 mmol) in DMF (50 mL) at room temperature was treated with Cs₂CO₃ (5.13 g, 15.7 mmol), stirred for 16 hours, poured into water, and extracted with ethyl acetate. The combined organics were dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 98:2 hexanes/ethyl acetate to provide 2.4 g (48%) of the desired product.

Example 95Bethyl 2-(5-((1,1'-biphenyl)-4-yloxy)pentyl)-1,3-dithiane-2-carboxylate

A suspension of NaH (224 mg, 8.9 mmol) in toluene (7 mL) at 0 °C was treated sequentially with ethyl-2-dithiane carboxylate (1.03 mL, 6.53 mmol) and a solution of Example 95A (2.3 g, 7.2 mmol) in DMF (2 mL), warmed to room temperature, stirred for 16 hours, poured into water, and extracted with ethyl acetate. The combined extracts were dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 98:2 hexanes/ethyl acetate to provide 0.52g (19%) of the desired product.

Example 95Cethyl 7-((3-bromo(1,1'-biphenyl)-4-yl)oxy)-2-oxoheptanoate

A solution of NBS (1.92g, 10.8 mmol) in a 97:3 mixture of acetone/water (19 mL) at 0 °C was treated dropwise with a solution of Example 95B (0.52g, 1.2 mmol) in a 97:3 mixture of acetone/water (3 mL), stirred for 15 minutes, and partitioned between

dichloromethane and 10% Na_2SO_3 . The organic phase was washed sequentially with 10% Na_2SO_3 , water, saturated NaHCO_3 , and brine, dried (Na_2SO_4), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with dichloromethane to provide 0.326g (65%) of the desired product. MS (ESI(+)) m/e 419 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 7.85 (d, 1H), 7.65-7.61 (m, 3H), 7.47-7.41 (m, 2H), 7.18 (d, 2H), 4.22 (q, 2H), 4.09 (t, 2H), 2.86 (t, 2H), 1.79-1.77 (m, 2H), 1.63-1.55 (m, 2H), 1.52-1.45 (m, 2H), 1.26 (t, 3H); Anal. Calcd. for $\text{C}_{21}\text{H}_{23}\text{BrO}_4 \cdot 0.5\text{H}_2\text{O}$: C, 58.89; H, 5.65. Found: C, 58.73; H, 5.37.

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Example 968-(1,3-oxazol-2-yl)-8-oxo-N-phenyloctanamide

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Example 96Amethyl 8-chloro-8-oxooctanoate

The desired product was prepared by substituting 8-methoxy-8-oxooctanoic acid for Example 102B in Example 102C.

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Example 96Bmethyl 8-(1,3-oxazol-2-yl)-8-oxooctanoate

The desired product was prepared by substituting Example 96A for Example 92C in Example 92D.

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Example 96C8-(1,3-oxazol-2-yl)-8-oxooctanoic acid

The desired product was prepared by substituting Example 96B for Example 1A in Example 1B.

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Example 96D8-(1,3-oxazol-2-yl)-8-oxo-N-phenyloctanamide

The desired product was prepared by substituting Example 96C for Example 1B in Example 1C. MS (ESI(+)) m/e 301 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.80 (s, 1H), 8.40 (s, 1H), 7.60 (d, 2H), 7.50 (s, 1H), 7.32 (t, 2H), 7.00 (t, 1H), 3.00 (t, 2H), 2.28 (t, 2H), 1.60 (m, 4H), 1.32 (m, 4H).

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Example 97N-(1,1'-biphenyl)-3-yl-8-(1,3-oxazol-2-yl)-8-oxooctanamide

The desired product was prepared by substituting Example 96C and (1,1'-biphenyl)-3-

amine for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+)) m/e 377 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.90 (s, 1H), 8.40 (s, 1H), 7.90 (s, 1H), 7.60 (dd, 2H), 7.50 (s, 1H), 7.48 (t, 2H), 7.30 (m, 4H), 3.00 (t, 2H), 2.28 (t, 2H), 1.60 (m, 4H), 1.32 (m, 4H).

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Example 98

N-(4-chlorophenyl)-8-(1,3-oxazol-2-yl)-8-oxooctanamide

The desired product was prepared by substituting Example 96C and 4-chloroaniline for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+)) m/e 335 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 10.00 (s, 1H), 8.40 (s, 1H), 7.60 (d, 2H), 7.50 (s, 1H), 7.30 (d, 2H), 3.00 (t, 2H), 2.28 (t, 2H), 1.60 (m, 4H), 1.32 (m, 4H).

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Example 99

8-(1,3-oxazol-2-yl)-8-oxo-N-(4-phenoxyphenyl)octanamide

The desired product was prepared by substituting Example 96C and 4-phenoxyaniline for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+)) m/e 393 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.82 (s, 1H), 8.40 (s, 1H), 7.60 (d, 2H), 7.50 (s, 1H), 7.40 (m, 2H), 7.10 (t, 1H), 7.00 (m, 4H), 3.00 (t, 2H), 2.28 (t, 2H), 1.60 (m, 4H), 1.34 (m, 4H).

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Example 100

8-(1,3-oxazol-2-yl)-8-oxo-N-(2-pyridinyl)octanamide

The desired product was prepared by substituting Example 96C and 2-aminopyridine for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+)) m/e 302 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 10.50 (s, 1H), 8.40 (s, 1H), 8.30 (m, 1H), 8.00 (d, 1H), 7.80 (m, 1H), 7.50 (s, 1H), 7.10 (m, 1H), 3.00 (t, 2H), 2.32 (t, 2H), 1.60 (m, 4H), 1.34 (m, 4H).

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Example 101

8-((1,1'-biphenyl)-4-yloxy)-N-methyl-2-oxooctanamide

The desired product was prepared by substituting methyl 7-bromoheptanoate for ethyl 7-bromoheptanoate in Example 2A.

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Example 101A

methyl 7-((1,1'-biphenyl)-4-yloxy)heptanoate

methyl 7-((1,1'-biphenyl)-4-yloxy)-2-hydroxyheptanoate

The desired product was prepared by substituting Example 101A for Example 2A in Example 94A.

5 Example 101C

8-((1,1'-biphenyl)-4-yloxy)-2-hydroxy-N-methyloctanamide

A suspension of Example 101B (20 mg, 0.06 mmol) and 2M dimethylamine in methanol (0.3 mL, 0.60 mmol) at room temperature was stirred for 48 hours and concentrated to provide 18 mg of the desired product. MS (ESI(+)) m/e 342 (M+H)⁺.

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Example 101D

8-((1,1'-biphenyl)-4-yloxy)-N-methyl-2-oxooctanamide

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The desired product was prepared by substituting Example 101A for Example 86A in Example 86B. MS (ESI(+)) m/e 340 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 7.62-7.56 (m, 4H), 7.45-7.40 (m, 2H), 7.33-7.27 (m, 1H), 7.03-6.98 (m, 2H), 4.00 (t, 2H), 2.82 (t, 2H), 2.64 (d, 3H), 1.77-1.68 (m, 2H), 1.58-1.30 (m, 6H); Anal. Calcd. for C₂₁H₂₅NO₃: C, 74.31; H, 7.42; N, 4.13. Found: C, 74.15; H, 7.60; N, 3.97.

20 Example 102

1-(1,3-oxazol-2-yl)-7-(phenylsulfanyl)-1-heptanone

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Example 102A

ethyl 7-(phenylsulfanyl)heptanoate

The desired product was prepared by substituting thiophenol for (1,1'-biphenyl)-4-ol in Example 2A.

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Example 102B

7-(phenylsulfanyl)heptanoic acid

The desired product was prepared by substituting Example 102A for Example 1A in Example 1B.

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Example 102C

7-(phenylsulfanyl)heptanoyl chloride

A solution of Example 102B (476 mg, 2 mmol) in dichloromethane at room temperature was treated with oxalyl chloride (0.26 mL, 3 mmol), stirred for 3 hours, and concentrated to provide the desired product.

Example 102D

1-(1,3-oxazol-2-yl)-7-(phenylsulfanyl)-1-heptanone

The desired product was prepared by substituting Example 102C for Example 92C in Example 92D. mp: 39-40 °C; MS (ESI(+)) m/e 290 (M+H)⁺; ¹H NMR (CDCl₃): 7.82 (m, 1H), 7.40-7.10 (m, 6H), 3.08 (t, 2H), 2.93 (t, 2H), 1.90-1.30 (m, 8H); Anal. Calcd. for C₁₆H₁₉NO₂S: C, 66.40; H, 6.62, N, 4.61. Found: C, 66.06; H, 6.31; N, 4.61.

Example 1031-(1,3-oxazol-2-yl)-7-(phenylsulfonyl)-1-heptanone

A solution of Example 102D (230 mg, 0.8 mmol) in 2:1 methanol:water (20 mL) at 10 room temperature was treated with Oxone[®] (1.22 g, 2 mmol) and NaHCO₃ (0.168 g, 2 mmol), stirred for 2 hours, and concentrated. The concentrate was partitioned between water and diethyl ether and the organic extract was washed with brine, dried (Na₂SO₄), filtered, and concentrated to provide 180 mg (70%) of the desired product. mp: 55-56 °C; MS (ESI(+)) m/e 322 (M+H)⁺; ¹H NMR (DMSO-d₆): 8.39 (s, 1H), 8.00-7.60 (m, 5H), 7.52 (s, 1H), 3.30 (m, 15 2H, overlap with H₂O), 2.98 (t, 2H), 1.70-1.20 (m, 8H); Anal. Calcd. for C₁₆H₁₉NO₄S: C, 59.79; H, 5.96; N 4.36. Found: C, 59.51; H, 6.13; N, 4.10.

Example 1047-(2-naphthylsulfanyl)-1-(1,3-oxazol-2-yl)-1-heptanone

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Example 104Aethyl 7-(2-naphthylsulfanyl)heptanoate

The desired product was prepared by substituting 2-naphthalenethiol for (1,1'-biphenyl)-4-ol in Example 2A.

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Example 104B7-(2-naphthylsulfanyl)heptanoic acid

The desired product was prepared by substituting Example 104A for Example 1A in Example 1B.

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Example 104C7-(2-naphthylsulfanyl)heptanoyl chloride

The desired product was prepared by substituting Example 104B for Example 102B in Example 102C.

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Example 104D1-(1,3-oxazol-2-yl)-7-(phenylsulfanyl)-1-heptanone

The desired product was prepared by substituting Example 104C for Example 92C in Example 92D. mp. 65-66 °C; MS (ESI(+)) m/e 340 (M+H)⁺; ¹H NMR (DMSO-d₆): 8.38 (s, 1H), 7.90-7.70 (m, 4H), 7.55-7.45 (m, 4H), 3.15-2.95 (m, 4H), 1.70-1.30 (m, 8H); Anal. Calcd. for C₂₀H₂₁NO₂S: C, 70.77; H, 6.24; N, 4.13. Found: C, 70.78; H, 6.47, N, 3.87.

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Example 105

7-(2-naphthylsulfonyl)-1-(1,3-oxazol-2-yl)-1-heptanone

The desired product was prepared by substituting Example 104D for Example 102D in Example 103. mp. 75-76 °C; MS (ESI(+)) m/e 372 (M+H)⁺; ¹H NMR (DMSO-d₆): 8.60-7.50 (m, 9H), 3.35 (m, 2H, overlap with H₂O), 2.98 (t, 2H), 1.65-1.20 (m, 8H); Anal. Calcd. for C₂₀H₂₁NO₄S: C, 64.67; H, 5.70; N, 3.77. Found: C, 64.39; H, 5.89; N, 3.53.

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Example 106

N-methyl-8-(2-naphthylsulfanyl)-2-oxooctanamide

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Example 106A

methyl 8-(2-naphthylsulfanyl)-2-oxooctanoate

The desired product was prepared by substituting 2-naphthalenethiol for 4-phenylphenol in Example 91.

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Example 106B

N-methyl-8-(2-naphthylsulfanyl)-2-oxooctanamide

A solution of Example 106A (0.8 g, 2.4 mmol) in THF (5 mL) at room temperature was treated with 2M methylamine in THF (2.4 mL, 4.8 mmol) and triethylamine (7 mL), 25 stirred for 4 hours, and concentrated. Recrystallization from ethyl acetate/hexanes provided 0.55g (69%) of the desired product. mp: 94-95 °C; MS (ESI(-)) m/e 328 (M-H)⁻; ¹H NMR (DMSO-d₆) δ 8.50 (br s, 1H), 7.90-7.30 (m, 7H), 3.07 (t, 2H), 2.79 (t, 2H), 2.64 (d, 3H), 1.70-1.20 (m, 8H); Anal. Calcd. for C₁₉H₂₃NO₂S: C, 69.27; H, 7.04; N, 4.25. Found: C, 68.92; H, 6.93; N, 4.05.

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Example 107

N-methyl-8-(2-naphthylsulfonyl)-2-oxooctanamide

The desired product was prepared by substituting Example 106B for Example 102D in Example 103. mp. 108-109 °C; MS (ESI(-)) m/e 360 (M-H)⁺; ¹H NMR (DMSO-d₆) δ 8.66 (s, 1H), 8.46 (br s, 1H), 8.30-7.60 (m, 6H), 3.35 (m, 2H, overlap with H₂O), 2.74 (t, 2H), 2.62 (d, 3H), 1.70-1.10 (m, 8H); Anal. Calcd. for C₁₉H₂₃NO₄S: C, 63.13; H, 6.41; N 3.88. Found: C, 62.93; H, 6.26; N, 3.53.

Example 1088-((1,1'-biphenyl)-4-ylsulfanyl)-N-methyl-2-oxooctanamide

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Example 108Amethyl 8-((1,1'-biphenyl)-4-ylsulfanyl)-2-oxooctanoate

The desired product was prepared by substituting (1,1'-biphenyl)-4-thiol for (1,1'-biphenyl)-4-ol in Example 91.

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Example 108B8-((1,1'-biphenyl)-4-ylsulfanyl)-N-methyl-2-oxooctanamide

The desired product was prepared by substituting Example 108A for Example 106A in Example 106B. mp: 131-132 °C; MS (ESI(-)) m/e 354 (M-H)⁻;

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¹H NMR (DMSO-d₆) δ 8.50 (br s, 1H), 7.70-7.30 (m, 9H), 3.00 (t, 2H), 2.79 (t, 2H), 2.62 (d, 3H), 1.70-1.20 (m, 8H); Anal. Calcd. for C₂₁H₂₅NO₂S: C, 70.95; H, 7.09; N, 3.94. Found: C 71.05, H 7.13, N, 3.79.

Example 1098-((1,1'-biphenyl)-4-ylsulfonyl)-N-methyl-2-oxooctanamide

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The desired product was prepared by substituting Example 108B for Example 102D in Example 103. mp: 134-135 °C; MS (ESI(-)) m/e 388 (M-H)⁻; ¹H NMR (DMSO-d₆) δ 8.50(br s, 1H), 7.80-7.40 (m, 9H), 3.40 (m, 2H, overlap with H₂O), 2.75 (t, 2H), 2.62(d, 3H), 1.60-1.20 (m, 8H).

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Example 110N-(7-(1,3-oxazol-2-yl)-7-oxoheptyl)-1H-indole-2-carboxamideExample 110Amethyl 7-((1H-indol-2-ylcarbonyl)amino)heptanoate

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The desired product was prepared by substituting methyl 7-aminoheptanoate and 1H-indole-2-carboxylic acid for Example 1B and aniline, respectively, in Example 1C.

Example 110B7-((1H-indol-2-ylcarbonyl)amino)heptanoic acid

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The desired product was prepared by substituting Example 110A for Example 1A in Example 1B.

Example 110C7-((1H-indol-2-ylcarbonyl)amino)heptanoyl chloride

The desired product was prepared by substituting Example 110B for Example 102B in Example 102C.

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Example 110DN-(7-(1,3-oxazol-2-yl)-7-oxoheptyl)-1H-indole-2-carboxamide

The desired product was prepared by substituting Example 110C for Example 92C in Example 92D. mp: 153-156 °C; MS (ESI(+)) m/e 340 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 1.40-10 1.30 (m, 4H), 1.70-1.50 (m, 4H), 3.06-3.01 (t, 2H), 3.34-3.24 (m, 2H), 7.04-6.99 (t, 1H), 7.09-7.08 (d, 1H), 7.18-7.13 (t, 1H), 7.43-7.40 (d, 1H), 7.52 (s, 1H), 7.60-7.58 (d, 1H), 8.38 (s, 1H), 8.44-8.40 (t, 1H), 11.51 (s, 1H); Anal. Calcd. for: C₁₉H₂₁N₃O₃: C, 67.24; H, 6.24; N, 12.38. Found: C, 66.98; H, 6.13; N, 12.05.

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Example 111N-(6-(1,3-oxazol-2-yl)-6-oxohexyl)-1H-indole-2-carboxamideExample 111Amethyl 6-((1H-indol-2-ylcarbonyl)amino)hexanoate

20 The desired product was prepared by substituting methyl 6-aminohexanoate and 1H-indole-2-carboxylic acid for Example 1B and aniline, respectively, in Example 1C.

Example 111B6-((1H-indol-2-ylcarbonyl)amino)hexanoic acid

25 The desired product was prepared by substituting Example 111A for Example 1A in Example 1B.

Example 111C6-((1H-indol-2-ylcarbonyl)amino)hexanoyl chloride

30 The desired product was prepared by substituting Example 111B for Example 102B in Example 102C.

Example 111DN-(6-(1,3-oxazol-2-yl)-6-oxohexyl)-1H-indole-2-carboxamide

35 The desired product was prepared by substituting Example 111C for Example 92C in Example 92D. mp: 176-179 °C; MS (ESI(+)) m/e 326 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 1.43-1.35 (m, 2H), 1.73-1.52 (m, 4H), 3.07-3.03 (t, 2H), 7.04-6.99 (t, 1H), 7.08 (s, 1H), 7.19-7.14

(t, 1H), 7.43-7.40 (d, 1H), 7.52 (s, 1H), 7.61-7.58 (d, 1H), 8.38 (s, 1H), 8.44-8.40 (t, 1H), 11.51 (s, 1H); Anal. Calcd. for $C_{18}H_{19}N_3O_3 \cdot 0.25H_2O$: C, 65.54; H, 5.95; N, 12.73. Found: C, 65.63; H, 5.78; N, 12.88.

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Example 1127-((1,1'-biphenyl)-4-yloxy)-1-(4,5-dihydro-1,3-oxazol-2-yl)-1-heptanoneExample 112A7-(1,1'-biphenyl-4-yloxy)heptan-1-ol

10 The desired product was prepared by substituting 7-(t-butyldimethylsilyloxy)heptyl bromide for 6-(t-butyldimethylsilyloxy)hexyl bromide in Example 91A.

Example 112B7-((1,1'-biphenyl)-4-yloxy)heptanal

15 The desired product was prepared by substituting Example 112A for Example 91A in Example 91B.

Example 112C8-((1,1'-biphenyl)-4-yloxy)-2-hydroxyoctanenitrile

20 A mixture of Example 112B (2.0g, 7.1 mmol) and KCN (4.66 g) in THF (25 mL) and water (27 mL) at room temperature was stirred for 2 days and concentrated. The resulting aqueous suspension was filtered to provide the desired product. MS (ESI(+)) m/e 327 ($M+NH_4$)⁺.

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Example 112D7-((1,1'-biphenyl)-4-yloxy)-1-(4,5-dihydro-1,3-oxazol-2-yl)-1-heptanol

A solution of acetyl chloride (2.86 mL) in ethanol (2.8 mL) and $CHCl_3$ (5.7 mL) at room temperature was treated with a solution of Example 112C (0.76 g, 2.5 mmol) in $CHCl_3$ (8 mL), stirred overnight, and concentrated. The concentrate was suspended in dichloromethane (12 mL), treated with ethanolamine (0.3 mL) and Et_3N (0.68 mL), stirred for 24 hours, and concentrated. The concentrate was suspended in $CHCl_3$ (50 mL), treated with $TsOH \cdot H_2O$ (46 mg), heated to reflux, and stirred for 3 hours. The reaction was cooled to room temperature, diluted with ethyl acetate, washed sequentially with saturated $NaHCO_3$, water, and brine, dried (Na_2SO_4), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 5% methanol/dichloromethane to provide 0.1 g of the desired product. MS (ESI(+)) m/e 354 ($M+H$)⁺.

Example 112E7-((1,1'-biphenyl)-4-yloxy)-1-(4,5-dihydro-1,3-oxazol-2-yl)-1-heptanone

The desired product was prepared by substituting Example 112D for Example 86A in Example 86B. MS (ESI(+)) m/e 352 (M+H)⁺; ¹H NMR (CDCl₃) δ 7.56-7.49 (m, 4H), 7.43-5 7.38 (m, 2H), 7.32-7.27 (m, 1H), 6.98-6.93 (m, 2H), 4.42 (t, 2H), 4.08 (t, 2H), 3.99 (t, 2H), 2.94 (t, 2H), 1.85-1.66 (m, 4H), 1.57-1.39 (m, 4H). Anal. Calcd. for C₂₂H₂₅NO₃: C, 75.19; H, 7.17; N, 3.99. Found: C, 75.32; H, 6.98; N, 3.89.

Example 11310 9-((1,1'-biphenyl)-4-yloxy)-2,3-nonanedioneExample 113A4-((5-iodopentyl)oxy)-1,1'-biphenyl

A suspension of 5-((4'-phenyl)phenoxy))-pentan-1-ol (0.85 g, 3.3 mmol) in dichloromethane (20 mL) was treated with methanesulfonyl chloride (0.28 mL, 3.6 mmol) and Et₃N (0.7 mL, 5 mmol), stirred at 0 °C for 1 hour, and partitioned between water and dichloromethane. The organic extract was washed with water, dried (Na₂SO₄), filtered, and concentrated. The concentrate was dissolved in acetone (30 mL), treated with NaI (2.5 g, 16.6 mmol), heated to 70 °C for 18 hours, cooled to room temperature, diluted with water, 15 and extracted three times with ethyl acetate. The combined organic extracts were washed sequentially with aqueous NaS₂O₃, water, and brine, dried (MgSO₄), filtered, and concentrated to provide 1.2 g of the desired product. MS (ESI(+)) m/e 384 (M+NH₄)⁺.
20

Example 113B25 methyl 7-((1,1'-biphenyl)-4-yloxy)-2-(2,2-dimethoxypropanoyl)heptanoate

A solution of NaH (64 mg, 1.6 mmol) in DMF (3 mL) at 0 °C was treated dropwise with methyl 4,4-dimethoxy-3-oxopentanoate (0.29 mL, 1.7 mmol), stirred for 1 hour, treated with a solution of Example 113A (0.5 g, 1.37 mmol) in DMF (4 mL), warmed to room temperature, and stirred for 36 hours. The reaction was quenched with water, and extracted 30 3 times with ethyl acetate. The combined organic extracts were washed with water and brine, dried (MgSO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 15% ethyl acetate/hexanes to provide 0.46 g (79%) of the desired product.

Example 113C35 9-((1,1'-biphenyl)-4-yloxy)-2,2-dimethoxy-3-nonanone

A solution of Example 113B (134 mg, 0.31 mmol) in methanol (4 mL) at room temperature was treated with 2N NaOH (0.63 mL), stirred for 3 hours, heated to 70 °C, stirred for 2 hours, cooled to room temperature, diluted with water, and extracted 3 times with ethyl acetate. The combined organic extracts were washed with water and brine, dried (MgSO₄), filtered, and concentrated to provide 92 mg (80%) of the desired product. mp: 48 °C.

Example 113D

9-((1,1'-biphenyl)-4-yloxy)-2,3-nonanedione

A solution of Example 113C (67 mg, 0.18 mmol) in THF (2 mL) and 4N HCl (1 mL) was heated at 40 °C for 1.5h. The reaction was cooled to r.t., diluted with water, extracted 2 times with EtOAc. The combined organic extracts were washed with water, brine, dried (MgSO₄), concentrated. The residue was triturated with hexane to give 43 mg (73% yield) of the title compound. mp: 110-111 °C; MS (Cl(+)) m/e 342 (M+NH₄)⁺; ¹H NMR (DMSO-d₆) δ 7.60 (m, 4H), 7.43 (t, 2H), 7.30 (t, 1H), 7.01 (d, 2H), 4.00 (t, 2H), 2.70 (t, 2H), 2.23 (s, 3H), 1.72 (m, 2H), 1.51 (quint, 2H), 1.48-1.30 (m, 4H); Anal. Calcd. for C₂₁H₂₄O₃: C, 77.75; H, 7.46. Found: C, 77.52; H, 7.19.

Example 114

N-(1,1'-biphenyl)-4-yl-7,8-dioxononanamide

Example 114A

6-ethyl 1-methyl 2-(2,2-dimethoxypropanoyl)hexanedioate

The desired product was prepared by substituting ethyl 4-bromobutanoate for Example 113A in Example 113B.

Example 114B

ethyl 7,7-dimethoxy-6-oxooctanoate

The desired product was prepared by substituting Example 114A for Example 113B in Example 113C.

Example 114C

N-(1,1'-biphenyl)-4-yl-7,7-dimethoxy-6-oxooctanamide

The desired product was prepared by substituting Example 114B and (1,1'-biphenyl)-4-amine for Example 1B and aniline, respectively, in Example 1C.

Example 114D

N-(1,1'-biphenyl)-4-yl-7,8-dioxononanamide

The desired product was prepared by substituting Example 114C for Example 113C in Example 113D. MS (Cl(+)) m/e 355 (M+NH₄)⁺; ¹H NMR (CDCl₃) δ 7.62-7.52 (m, 4H), 7.42 (t, 2H), 7.32 (t, 1H), 7.02 (br s, 1H), 2.77 (t, 2H), 2.39 (t, 2H), 2.33 (s, 3H), 1.77 (quint, 2H), 1.65 (quint, 2H), 1.48-1.36 (m, 2H).

Example 115methyl 7-((1,1'-biphenyl)-4-yloxy)-2-oxoheptanoate

The desired product was prepared by substituting 5-(t-butyldimethylsilyloxy)pentyl bromide (prepared according to the procedure described in *Can. J. Chem.* **1994**, 72, 1500-1511) for 6-(t-butyldimethylsilyloxy)hexyl bromide in Example 91. MS (ESI(+)) m/e 327 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 7.62-7.56 (m, 4H), 7.44-7.39 (t, 2H), 7.32-7.27 (m, 1H), 7.00 (d, 2H), 4.00 (t, 2H), 3.77 (s, 3H), 2.86 (t, 2H), 1.78-1.69 (m, 2H), 1.63-1.53 (m, 2H), 1.49-1.40 (m, 2H).

15

Example 116methyl 9-((1,1'-biphenyl)-3-ylamino)-2,9-dioxononanoateExample 116AN-(1,1'-biphenyl)-3-yl-6-(1,3-dioxolan-2-yl)hexanamide

The desired product was prepared by substituting 6-(1,3-dioxolan-2-yl)hexanoic acid and (1,1'-biphenyl)-3-amine for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+)) m/e 340 (M+H)⁺.

25

Example 116BN-(1,1'-biphenyl)-3-yl-7-oxoheptanamide

A solution of Example 116A (2.35g, 6.9 mmol) in acetone (20 mL) and water (2 mL) was treated with TsOH·H₂O (15 mg), heated to reflux, stirred overnight, cooled to room temperature, and concentrated. The concentrate was dissolved in ethyl acetate, washed with water, dried (Na₂SO₄), filtered, and concentrated to provide 1.82 g (89%) of the desired product. MS (ESI(+)) m/e 296 (M+H)⁺.

Example 116Cmethyl 9-((1,1'-biphenyl)-3-ylamino)-2,9-dioxononanoate

The desired product was prepared by substituting Example 116B for Example 91B in Example 91C. MS (ESI(+)) m/e 368 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.95 (s, 1H), 7.92 (br

s, 1H), 7.62-7.55 (m, 3H), 7.50-7.43 (m, 2H), 7.40-7.28 (m, 3H), 3.76 (s, 3H), 2.82 (t, 2H), 2.32 (t, 2H), 1.64-1.47 (m, 4H), 1.46-1.37 (m, 4H).

Example 117

5

methyl 9-anilino-2,9-dioxononanoate

Example 117A

6-hydroxy-N-phenylhexanamide

The desired product was prepared by substituting 6-hydroxyhexanoic acid for
10 Example 1B in Example 1C.

Example 117B

methyl 9-anilino-2,9-dioxononanoate

The desired product was prepared by substituting Example 117A for Example 91A in
15 Example 91. MS (ESI(+)) m/e 292 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.82 (s, 1H), 7.57 (d, 2H), 7.27 (t, 2H), 7.01 (t, 1H), 3.76 (s, 3H), 2.81 (t, 2H), 2.28 (t, 2H), 1.62-1.48 (m, 4H), 1.33-1.26 (m, 4H).

Example 118

20 methyl 8-((4'-cyano(1,1'-biphenyl)-4-yl)oxy)-2-oxooctanoate

The desired product was prepared by substituting 4'-hydroxy(1,1'-biphenyl)-4-carbonitrile for (1,1'-biphenyl)-4-ol in Example 91. MS (ESI(-)) m/e 364 (M-H)⁻.

Example 119

25 8-((4'-cyano(1,1'-biphenyl)-4-yl)oxy)-N-methyl-2-oxooctanamide

A suspension of Example 118 (261 mg, 0.7 mmol) and MeNH₂·HCl (150 mg, 2.2 mmol) in CH₃CN (5 mL) and Et₃N (10 mL) at room temperature was stirred in a sealed vessel for 18 hours, then partitioned between water and ethyl acetate. The organic extract was dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash 30 column chromatography on silica gel with 30% ethyl acetate/hexanes to provide 113 mg (44%) of the desired product. MS (ESI(-)) m/e 363 (M-H)⁻; ¹H NMR (DMSO-d₆) δ 8.53-8.48 (m, 1H), 7.90-7.83 (m, 4H), 7.70 (d, 2H), 7.05 (d, 2H), 4.04-4.00 (m, 2H), 2.82 (t, 2H), 2.64 (d, 3H), 1.78-1.68 (m, 2H), 1.58-1.47 (m, 2H), 1.47-1.30 (m, 4H).

Example 120

35 N⁹-(1,1'-biphenyl)-3-yl-N¹-methyl-2-oxononanediamide

The desired product was prepared by substituting Example 116 for Example 118 in Example 119. MS (ESI(+)) m/e 367 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.95 (s, 1H), 8.53-8.48 (m, 1H), 7.92-7.90 (m, 1H), 7.61-7.54 (m, 3H), 7.50-7.43 (m, 2H), 7.40-7.28 (m, 3H), 2.80 (t, 2H), 2.64 (d, 3H), 2.32 (t, 2H), 1.63-1.55 (m, 2H), 1.55-1.47 (m, 2H), 1.34-1.28 (m, 4H).

5

Example 121(7E)-8-(1,1'-biphenyl)-4-yl-1-(1,3-oxazol-2-yl)-7-octen-1-oneExample 121A(7E)-8-(1,1'-biphenyl)-4-yl-7-octenoic acid

10

The desired product was prepared by substituting Example 87A for Example 1A in Example 1B.

Example 121B(7E)-8-(1,1'-biphenyl)-4-yl-7-octenoyl chloride

15

The desired product was prepared by substituting Example 121A for Example 102B in Example 102C.

Example 121D(7E)-8-(1,1'-biphenyl)-4-yl-1-(1,3-oxazol-2-yl)-7-octen-1-one

20

The desired product was prepared by substituting Example 121C for Example 92C in Example 92D. MS (ESI(+)) m/e 346 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.39 (s, 1H), 7.70-7.64 (m, 4H), 7.52 (s, 1H), 7.50-7.43 (m, 2H), 7.40-7.32 (m, 3H), 6.44 (br d, 1H), 5.69 (dt, 1H), 3.03 (t, 2H), 2.40-2.31 (m, 2H), 1.69-1.59 (m, 2H), 1.53-1.32 (m, 4H).

25

Example 1224-((1E)-8-(1,3-oxazol-2-yl)-8-oxo-1-octenyl)benzonitrileExample 122Aethyl (7E)-8-(4-cyanophenyl)-7-octenoate

30

The desired product was prepared by substituting 4-cyanobenzaldehyde for 4-phenylbenzaldehyde in Example 87A.

Example 122B(7E)-8-(4-cyanophenyl)-7-octenoic acid

35

The desired product was prepared by substituting Example 122A for Example 1A in Example 1B.

Example 122C(7E)-8-(4-cyanophenyl)-7-octenoyl chloride

The desired product was prepared by substituting Example 122B for Example 102B in
5 Example 102C.

Example 122D4-((1E)-8-(1,3-oxazol-2-yl)-8-oxo-1-octenyl)benzonitrile

The desired product was prepared by substituting Example 122C for Example 92C in
10 Example 92D. MS (ESI(+)) m/e 295 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.38 (s, 1H), 7.80 (d, 2H), 7.53 (s, 1H), 7.46 (d, 2H), 6.48 (br d, 1H), 5.89-5.79 (m, 1H), 3.01 (t, 2H), 2.34-2.27 (m, 2H), 1.67-1.56 (m, 2H), 1.51-1.30 (m, 4H).

Example 123N⁹-(1,1'-biphenyl)-3-yl-2-oxanonanediame

A suspension of Example 116 (354mg, 0.96mmol) in ethanol (5 mL) at room
temperature was treated with concentrated NH₄OH (1 mL), stirred for 2 hours, and filtered.
The isolated solid was washed with ethanol and dried under vacuum with heating to provide
158 mg (47%) of the desired product. MS (ESI(+)) m/e 353 (M+H)⁺; ¹H NMR (DMSO-d₆) δ
15 9.96 (s, 1H), 7.94-7.89 (m, 2H), 7.64-7.54 (m, 4H), 7.50-7.43 (m, 2H), 7.41-7.29 (m, 3H),
20 2.78 (t, 2H), 2.32 (t, 2H), 1.63-1.54 (m, 2H), 1.54-1.45 (m, 2H), 1.34-1.26 (m, 4H).

Example 124N¹-methyl-2-oxo-N⁹-(4-phenyl-1,3-thiazol-2-yl)nonanediame

25
A solution of methyl 6-hydroxyhexanoate (5.0 g, 31.0 mmol, prepared according to
the procedure described in Syn.Comm. 1991, 21 1075) in THF (11 mL) was treated with 2M
30 LiOH (16 mL), heated to 60 °C for 4 hours, heated to 85 °C for 2 hours, and concentrated
under nitrogen. The crude product was suspended in DMF (100 mL), treated with
NaHCO₃ (5.26 g, 6.3 mmol) and a solution of benzyl bromide (9.3 mL, 78 mmol) in DMF (50
mL), stirred for 18 hours, poured into water, and extracted three times with ethyl acetate.
The combined extracts were dried (Na₂SO₄), filtered, and concentrated. The concentrate was
35 purified by flash column chromatography on silica gel with 10% ethyl acetate/hexanes to
30% ethyl acetate/hexanes to provide 4.43g (60%) of the desired product. MS (ESI(+)) m/e
237 (M+H)⁺.

Example 124B
benzyl 7-oxoheptanoate

The desired product was prepared by substituting Example 124A for Example 91A in
5 Example 91B.

Example 124C
9-benzyl 1-methyl 2-oxanonanedioate

The desired product was prepared by substituting Example 124B for Example 91B in
10 Example 91C.

Example 124D
benzyl 9-(methylamino)-8,9-dioxanonanoate

The desired product was prepared by substituting Example 124C for Example 118 in
15 Example 119. MS (ESI(+)) m/e 306 (M+H)⁺.

Example 124E
9-(methylamino)-8,9-dioxanonanoic acid

A solution of Example 124D (0.89 g, 2.91 mmol) and 10% Pd/C (95 mg) in methanol
20 (15 mL) was stirred for 1 hour under a hydrogen atmosphere, filtered through diatomaceous
earth (Celite[®]), and concentrated to provide 0.56g (89 %) of the desired product. MS
(ESI(+)) m/e 216 (M+H)⁺.

Example 124F
N¹-methyl-2-oxo-N⁹-(4-phenyl-1,3-thiazol-2-yl)nonanediamide

The desired product was prepared by substituting Example 124E and 4-phenyl-1,3-thiazol-2-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (APCI(+)) m/e 374 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.20 (s, 1H), 8.53-8.74 (br m, 1H), 7.89 (d, 2H), 7.59 (s, 1H), 7.43 (t, 2H), 7.34-7.29 (m, 1H), 2.80 (t, 2H), 2.64 (d, 3H), 2.44 (t, 2H), 1.65-1.55 (m, 2H), 1.55-1.45 (m, 2H), 1.32-1.26 (m, 4H).

Example 125
N¹-methyl-2-oxo-N⁹-(4-phenoxyphenyl)nonanediamide

The desired product was prepared by substituting Example 124E and 4-phenoxyaniline for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 383 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.87 (s, 1H), 8.52-8.47 (br m, 1H), 7.59

(d, 2H), 7.48-7.43 (m, 2H), 7.11-7.06 (m, 1H), 6.98-7.93 (m, 4H), 2.80 (t, 2H), 2.64 (d, 3H), 2.28 (t, 2H), 1.62-1.45 (m, 4H), 1.32-1.26 (m, 4H).

Example 126

5 N^9 -(4,5-diphenyl-1,3-thiazol-2-yl)- N^1 -methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4,5-diphenyl-1,3-thiazol-2-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 450 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.26 (s, 1H), 8.52-8.47 (br m, 1H), 7.44-7.29 (m, 10H), 2.80 (t, 2H), 2.64 (d, 3H), 2.45 (t, 2H), 1.64-1.56 (m, 2H), 1.54-1.44 (m, 2H), 10 1.33-1.26 (m, 4H).

Example 127

N^9 -(4-(3-methoxyphenyl)-1,3-thiazol-2-yl)- N^1 -methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-(3-methoxyphenyl)-1,3-thiazol-2-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 404 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.19 (s, 1H), 8.53-8.47 (br m, 1H), 7.62 (s, 1H), 7.49-7.44 (m, 2H), 7.38-7.27 (m, 1H), 6.91-6.84 (m, 1H), 3.80 (s, 3H), 2.80 (t, 2H), 2.64 (d, 3H), 2.44 (t, 2H), 1.64-1.55 (m, 2H), 1.53-1.43 (m, 2H), 1.32-1.23 (m, 4H).

20

Example 128

N^9 -(4-(2-methoxyphenyl)-1,3-thiazol-2-yl)- N^1 -methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-(2-methoxyphenyl)-1,3-thiazol-2-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 404 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.12 (s, 1H), 8.53-8.47 (br m, 1H), 8.05 (dd, 1H), 7.62 (s, 1H), 7.33-7.37 (m, 1H), 7.14-7.11 (m, 1H), 7.05-7.00 (m, 1H), 3.91 (s, 3H), 2.80 (t, 2H), 2.64 (d, 3H), 2.44 (t, 2H), 1.64-1.56 (m, 2H), 1.56-1.44 (m, 2H), 1.32-1.25 (m, 4H).

30

Example 129

N^1 -methyl- N^9 -(5-methyl-4-phenyl-1,3-thiazol-2-yl)-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 5-methyl-4-phenyl-1,3-thiazol-2-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 388 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.03 (s, 1H), 8.52-8.47 (br m, 1H), 7.54-7.51 (m, 2H), 7.44 (t, 2H), 7.34-7.31 (m, 1H), 2.79 (t, 2H), 2.64 (d, 3H), 2.46 (s, 3H), 2.41 (t, 2H), 1.62-1.54 (m, 2H), 1.54-1.44 (m, 2H), 1.30-1.25 (m, 4H).

Example 130N¹-methyl-2-oxo-N⁹-(4'-(trifluoromethoxy)(1,1'-biphenyl)-3-yl)nonanediamide

The desired product was prepared by substituting Example 124E and 4-(4-trifluoromethoxyphenoxy)aniline for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 451 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.98 (s, 1H), 8.53-8.48 (br m, 1H), 8.05-8.03 (m, 1H), 7.72 (d, 2H), 7.59-7.56 (m, 1H), 7.48-7.30 (m, 4H), 2.80 (t, 2H), 2.64 (d, 3H), 2.32 (t, 2H), 1.64-1.55 (m, 2H), 1.55-1.46 (m, 2H), 1.33-1.28 (m, 4H).

Example 131N⁹-(4-(4-chlorophenoxy)phenyl)-N¹-methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-(4-chlorophenoxy)aniline for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 417 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.89 (s, 1H), 8.53-8.46 (br m, 1H), 7.61 (d, 2H), 7.40 (d, 2H), 7.01-6.95 (m, 4H), 2.80 (t, 2H), 2.64 (d, 3H), 2.28 (t, 2H), 1.62-1.45 (m, 4H), 1.33-1.26 (m, 4H).

Example 132N⁹-(4-methoxy(1,1'-biphenyl)-3-yl)-N¹-methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-methoxy(1,1'-biphenyl)-3-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 397 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.11 (s, 1H), 8.53-8.47 (br m, 1H), 8.30-8.28 (br s, 1H), 7.56 (d, 2H), 7.46-7.41 (m, 2H), 7.38-7.29 (m, 2H), 7.12 (d, 1H), 3.87 (s, 3H), 2.81 (t, 2H), 2.64 (d, 3H), 2.43-2.36 (m, 2H), 1.63-1.48 (m, 4H), 1.34-1.27 (m, 4H).

Example 133N⁹-(4'-cyano(1,1'-biphenyl)-3-yl)-N¹-methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 3'-amino(1,1'-biphenyl)-4-carbonitrile for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 392 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 10.01 (s, 1H), 8.52-8.47 (br m, 1H), 8.02-8.00 (br m, 1H), 7.93 (d, 2H), 7.80 (d, 2H), 7.63-7.59 (m, 1H), 7.46-7.38 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.32 (t, 2H), 1.64-1.45 (m, 4H), 1.33-1.28 (m, 4H).

Example 134N⁹-(4-bromophenyl)-N¹-methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-bromoaniline for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 368

(M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.97 (s, 1H), 8.52-8.47 (br s, 1H), 7.55 (d, 2H), 7.46 (d, 2H), 2.79 (t, 2H), 2.64 (d, 3H), 2.28 (t, 2H), 1.61-1.44 (m, 4H), 1.32-1.24 (m, 4H).

Example 135

5 N⁹-(6-methoxy(1,1'-biphenyl)-3-yl)-N¹-methyl-2-oxanonanediamide

The desired product was prepared by substituting Example 124E and 6-methoxy(1,1'-biphenyl)-3-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 397 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.76 (s, 1H), 8.52-8.47 (br m, 1H), 7.54-7.51 (m, 2H), 7.45-7.36 (m, 4H), 7.34-7.29 (m, 1H), 7.03 (d, 1H), 3.72 (s, 3H), 2.79 (t, 2H), 10 2.64 (d, 3H), 2.26 (t, 2H), 1.61-1.45 (m, 4H), 1.32-1.25 (m, 4H).

Example 136

N⁹-(1,1'-biphenyl)-4-yl-N¹-methyl-2-oxanonanediamide

The desired product was prepared by substituting Example 124E and (1,1'-biphenyl)-4-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 367 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.94 (s, 1H), 8.52-8.47 (br m, 1H), 7.70-7.58 (m, 6H), 7.44 (t, 2H), 7.32 (t, 1H), 2.80 (t, 2H), 2.64 (d, 3H), 2.32 (t, 2H), 1.63-1.56 (m, 2H), 1.55-1.46 (m, 2H), 1.33-1.27 (m, 4H).

20 Example 137

N⁹-(3,4-dichlorophenyl)-N¹-methyl-2-oxanonanediamide

The desired product was prepared by substituting Example 124E and 3,4-dichloroaniline for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 359 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 10.15 (s, 1H), 8.53-8.47 (br s, 1H), 7.99 (d, 1H), 7.54 (d, 1H), 7.47 (dd, 1H), 2.79 (t, 2H), 2.64 (d, 3H), 2.30 (t, 2H), 1.62-1.44 (m, 4H), 1.32-1.25 (m, 4H).

Example 138

N¹-methyl-2-oxo-N⁹-(4-(trifluoromethyl)phenyl)nonanediamide

30 The desired product was prepared by substituting Example 124E and 4-trifluoromethylaniline for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 359 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 10.21 (s, 1H), 8.52-8.47 (br m, 1H), 7.80 (d, 2H), 7.64 (d, 2H), 2.79 (t, 2H), 2.64 (d, 3H), 2.33 (t, 2H), 1.63-1.44 (m, 4H), 1.33-1.26 (m, 4H).

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Example 139

N⁹-(3-cyanophenyl)-N¹-methyl-2-oxanonanediamide

The desired product was prepared by substituting Example 124E and 3-cyanoaniline for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 316 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 10.20 (s, 1H), 8.53-8.46 (br m, 1H), 8.10-8.08 (br m, 1H), 7.80-7.76 (m, 1H), 7.51-7.47 (m, 2H), 2.79 (t, 2H), 2.64 (d, 3H), 2.32 (t, 2H), 1.63-1.44 (m, 4H), 1.32-1.26 (m, 4H).

5 **Example 140**

N⁹-(4-(4-methoxyphenyl)-1,3-thiazol-2-yl)-N¹-methyl-2-oxanonanediamide

The desired product was prepared by substituting Example 124E and 4-(4-methoxyphenyl)-1,3-thiazol-2-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 404 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.15 (s, 1H), 8.53-8.48 (br m, 1H), 7.81 (d, 2H), 7.42 (s, 1H), 6.98 (d, 2H), 3.79 (s, 3H), 2.80 (t, 2H), 2.64 (d, 3H), 2.43 (t, 2H), 1.64-1.54 (m, 2H), 1.54-1.44 (m, 4H), 1.31-1.26 (m, 4H).

15 **Example 141**

methyl 8-((4'cyano(1,1'-biphenyl)-4-yl)oxy)-3-hydroxy-2oxooctanoate

Example 141A

4'-((6-oxohexyl)oxy)-1,1'-biphenyl-4-carbonitrile

20 The desired product was prepared by substituting 4'-hydroxy-1,1'-biphenyl-4-carbonitrile for (1,1'-biphenyl)-4-ol in Examples 91A and 91B. MS (ESI(-)) m/e 292 (M-H)⁻.

Example 141B

methyl (2Z)-2-((tert-butyl(dimethyl)silyl)oxy)-8-((4'-cyano-1,1'-biphenyl-4-yl)oxy)oct-2-enoate

25 The desired product was prepared by substituting Example 141A for Example 142B in Example 142C. MS (ESI(+)) m/e 480 (M+H)⁺.

Example 141C

30 methyl 2-((tert-butyl(dimethyl)silyl)oxy)-3-((4'-cyano-1,1'-biphenyl-4-yl)oxy)pentyl oxirane-2-carboxylate

35 A solution of Example 141B (0.68 g, 1.42 mmol) in dichloromethane (10 mL) at room temperature was treated with 70% m-CPBA (350 mg, 1.42 mmol), stirred for 24 hours, treated with activated KF (150 mg), and stirred for 3 hours. The suspension was filtered and the filtrate was concentrated and purified by flash column chromatography on silica gel with 5:1 hexanes/ethyl acetate to provide 0.5 g (71% yield) of the desired product.

Example 141Dmethyl 8-((4'cyano(1,1'-biphenyl)-4-yl)oxy)-3-hydroxy-2oxooctanoate

An solution of Example 141C (50 mg, 0.1 mmol) in acetonitrile (1 mL) at 0 °C was treated with Et₃N·HF (2 drops), warmed to room temperature, stirred for 24 hours, adjusted to pH 7 with saturated NaHCO₃, diluted with water, and filtered. The filter cake was washed with water and dried under vacuum to provide 10 mg (26% yield) of the desired product. MS (DCI) m/e 399 (M+NH₄)⁺; ¹H NMR (DMSO-d₆) δ 7.89-7.82 (m, 4H), 7.72-7.68 (m, 2H), 7.06-7.03 (m, 2H), 6.84 (d, 1H), 4.22 (d, 1H), 4.00 (t, 2H), 3.70 (s, 3H), 1.74-1.64 (m, 2H), 1.44-1.18 (m, 6H); Anal Calcd for C₂₂H₂₃NO₅·0.25H₂O: C, 68.47; H, 6.14; N, 3.63. Found: C, 68.63; H, 6.21; N, 3.45.

Example 142N-(8-(methylamino)-7,8-dioxooctyl)-1H-indole-2-carboxamide15 Example 142Atert-butyl 6-hydroxyhexylcarbamate

A solution of 6-aminohexan-1-ol (5.03 g, 42.9 mmol) in THF (35 mL) at room temperature was treated portionwise with (Boc)₂O (8.9 g, 40.1 mmol), stirred for 1 hour, and filtered. The filtrate was concentrated, diluted with diethyl ether, washed sequentially with 1M HCl, water, and brine, dried (MgSO₄), and concentrated to provide the desired product (7.83 g). MS (ESI(+)) m/e 218 (M+H)⁺.

Example 142Btert-butyl 6-oxohexylcarbamate

25 A solution of oxalyl chloride (3.64 mL, 41.7 mmol) in dichloromethane (200 mL) at -78 °C was treated dropwise with DMSO (6 mL, 84.6 mmol), stirred for 5 minutes, treated with a solution of Example 142A (7.56 g, 34.8 mmol) in dichloromethane (100 mL), stirred for 15 minutes, treated with triethylamine (24 mL, 172 mmol), and warmed to room temperature. The reaction was partitioned between water and dichloromethane and the 30 organic phase was washed with brine, dried (MgSO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 10 to 20% ethyl acetate/hexanes to provide 6.44g (86% yield) of the desired product.

Example 142C35 methyl (2Z)-8-((tert-butoxycarbonyl)amino)-2-((tert-butyl(dimethyl)silyl)oxy)oct-2-enoate

A suspension of LiCl (1.5 g, 36 mmol) in THF (50 mL) at room temperature was treated with a solution of methyl ((tert-

butyl(dimethylsilyl)oxy(dimethoxyphosphoryl)acetate (6.25 g, 20 mmol) in THF (25 mL), treated with DBU (3.6 mL, 24 mmol), stirred for 15 minutes, cooled to 0 °C, treated with a solution of Example 142B (4.3 g, 20 mmol) in THF (25 mL), cooled to 0 °C, stirred for 30 minutes, warmed to room temperature, and stirred for 18 hours. The mixture was diluted 5 with ethyl acetate, washed with water and brine, dried (Na₂SO₄), filtered, and concentrated to provide the desired product (7.81 g). MS (ESI(+)) m/e 402 (M+H)⁺.

Example 142D

methyl 8-((tert-butoxycarbonyl)amino)-2-oxooctanoate

10 A solution of Example 142C (7.81 g, 19.4 mmol) in acetonitrile (200 mL) at 0 °C was treated with acetic acid (4 mL, 97 mmol) and CsF (5.89 g, 38 mol), stirred at 0 °C for 1 hour, warmed to room temperature, and stirred for 18 hours. The reaction mixture was diluted with 1:1 hexanes/ethyl acetate (400 mL), washed sequentially with NaHCO₃ (9.8 g in 200 mL water), water, and brine, dried (MgSO₄), filtered, and concentrated to provide 4.43g (79%) of 15 the desired product. MS (ESI(+)) m/e 286 (M+H)⁺.

Example 142E

tert-butyl 8-(methylamino)-7,8-dioxooctylcarbamate

20 The desired product was prepared by substituting Example 142D for Example 106A in Example 106B.

Example 142F

8-amino-N-methyl-2-oxooctanamide

25 Example 142E (1.06 g, 3.9 mmol) at room temperature was treated with 4N HCl in dioxane (10 mL), stirred for 1 hour, and concentrated under a stream of nitrogen to provide 0.809 g of the desired product as the hydrochloride salt. MS (ESI(+)) m/e 187 (M+H)⁺.

Example 142G

N-(8-(methylamino)-7,8-dioxooctyl)-1H-indole-2-carboxamide

30 The desired product was prepared by substituting Example 142F and 1H-indole-2-carboxylic acid for aniline and Example 1B, respectively, in Example 1C. MS (ESI(+)) m/e 330 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 11.50 (br s, 1H), 8.53-8.48 (br m, 1H), 8.43-8.39 (br m, 1H), 7.59 (d, 1H), 7.41 (d, 1H), 7.19-7.13 (m, 1H), 7.08 (d, 1H), 7.04-6.99 (m, 1H), 3.29-3.23 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.58-1.47 (m, 4H), 1.37-1.28 (m, 4H).

35

Example 143

N-(7-(methylamino)-6,7-dioxoheptyl)-1H-indole-2-carboxamide

Example 143A7-amino-N-methyl-2-oxoheptanamide

The desired product was prepared as the hydrochloride salt by substituting 5-
5 aminohexan-1-ol for 6-aminohexan-1-ol in Examples 142A-142F.

Example 143BN-(7-(methylamino)-6,7-dioxoheptyl)-1H-indole-2-carboxamide

The desired product was prepared by substituting Example 143A and 1H-indole-2-
10 carboxylic acid for aniline and Example 1B, respectively, in Example 1C. MS (ESI(+)) m/e 316 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 11.51 (s, 1H), 8.52-8.47 (br m, 1H), 8.43-8.39 (br m, 1H), 7.59 (d, 1H), 7.41 (d, 1H), 7.19-7.13 (m, 1H), 7.08 (d, 1H), 7.05-6.99 (m, 1H), 3.31-3.23 (m, 2H), 2.82 (t, 2H), 2.64 (d, 3H), 1.59-1.49 (m, 4H), 1.38-1.28 (m, 2H).

15

Example 144N-(7-(methylamino)-6,7-dioxoheptyl)-1,1'-biphenyl-4-carboxamide

Example 143A was coupled with 4-phenylbenzoic acid following the procedures of
Example 1C to provide the desired product. MS (ESI(+)) m/e 353 (M+H)⁺; ¹H NMR
(DMSO-d₆) δ 8.52-8.46 (m, 2H), 7.93 (d, 2H), 7.77-7.71 (m, 4H), 7.52-7.47 (m, 2H), 7.43-
20 7.40 (m, 1H), 3.31-3.25 (m, 2H), 2.82 (t, 2H), 2.64 (d, 3H), 1.59-1.49 (m, 4H), 1.37-1.29 (m, 2H).

Example 1457-((4-chlorophenyl)sulfonyl)amino-N-methyl-2-oxoheptanamide

25 A solution of Example 143A (125 mg, 0.6 mmol) in DMF (3 mL) at room
temperature was treated with 4-chlorophenylsulfonyl chloride (127 mg, 0.6 mmol) and Et₃N
(0.17 mL, 1.2 mmol), stirred for 18 hours, and treated with cold water. The precipitate was
collected by filtration and the filter cake was washed with water and dried under vacuum to
provide 156 mg (75%) of the desired product. MS (ESI(+)) m/e 347, 349 (M+H)⁺; ¹H NMR
(DMSO-d₆) δ 8.52-8.48 (br m, 1H), 7.79 (d, 2H), 7.68-7.65 (m, 3H), 2.77-2.69 (m, 4H), 2.64
30 (d, 3H), 1.46-1.31 (m, 4H), 1.25-1.17 (m, 2H).

Example 146N~1~-methyl-2-oxo-N~9~-phenylnonanediamide

35 The desired product was prepared by substituting Example 124E and aniline for
Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 291
(M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.82 (s, 1H), 8.53-8.47 (m, 1H), 7.59-7.56 (m,

2H),), 7.29 (t, 2H), 7.04-6.98 (m, 1H), 7.04-6.98 (m, 1H), 2.64 (d, 3H), 2.28 (t, 2H), 1.63-1.45 (m, 4H), 1.32-1.27 (m, 4H).

Example 147

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N~1~-methyl-2-oxo-N~9~-{(1,3-thiazol-2-yl)nonanediamide}

The desired product was prepared by substituting Example 124E and 2-aminothiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 298 (M+H)⁺;

10 ¹H NMR (300 MHz, DMSO-d₆) δ 12.02 (s, 1H), 8.53-8.48 (m, 1H), 7.44 (d, 1H), 7.17 (d, 1H), 2.79 (t, 2H), 2.64 (d, 3H), 2.41 (t, 2H), 1.63-1.45 (m, 4H), 1.30-1.24 (m, 4H).

Example 148

N~9~-{(4-methoxyphenyl)-N~1~-methyl-2-oxononanediamide}

15 The desired product was prepared by substituting Example 124E and 4-methoxyaniline for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 321 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.68 (s, 1H), 8.53-8.46 (m, 1H), 7.47 (d, 2H), 6.85 (d, 2H), 3.71 (s, 3H), 2.79 (t, 2H), 2.64 (d, 3H), 2.24 (t, 2H), 1.61-1.46 (m, 4H), 1.32-1.26 (m, 4H).

20

Example 149

N~9~-{(4-chlorophenyl)-N~1~-methyl-2-oxononanediamide}

15 The desired product was prepared by substituting Example 124E and 4-chloroaniline for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 325, 327 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.97 (s, 1H), 8.53-8.47 (m, 1H), 7.61 (d, 2H), 7.33 (d, 2H), 2.79 (t, 2H), 2.64 (d, 3H), 2.29 (t, 2H), 1.61-1.45 (m, 4H), 1.34-1.26 (m, 4H).

Example 150

N~1~-methyl-N~9~-{(2-naphthyl)-2-oxononanediamide}

20 The desired product was prepared by substituting Example 124E and 2-aminonaphthalene for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 341 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 10.05 (s, 1H), 8.55-8.47 (m, 1H), 8.29 (br s, 1H), 7.85-7.77 (m, 3H), 7.59-7.55 (m, 1H), 7.48-7.35 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.35 (t, 2H), 1.66-1.57 (m, 2H), 1.51-1.47 (m, 2H), 1.34-1.28 (m, 4H).

35

Example 151

N~1~-methyl-2-oxo-N~9~-{quinolin-3-yl}nonanediamide

The desired product was prepared by substituting Example 124E and 3-aminoquinoline for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 342 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 10.41 (s, 1H), 8.95 (d, 1H), 8.75 (d, 1H), 8.54-8.47 (m, 1H), 7.98-7.93 (m, 2H), 7.70-7.64 (m, 1H), 7.62-7.57 (m, 1H), 5 2.81 (t, 2H), 2.64 (d, 3H), 2.40 (t, 2H), 1.69-1.58 (m, 2H), 1.56-1.45 (m, 2H), 1.48-1.39 (m, 4H).

Example 152

N~9~-(1,3-benzothiazol-2-yl)-N~1~-methyl-2-oxononanediamide

10 The desired product was prepared by substituting Example 124E and 2-aminobenzothiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 348 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 12.28 (s, 1H), 8.54-8.48 (m, 1H), 7.97-7.94 (m, 1H), 7.74-7.71 (m, 1H), 7.45-7.40 (m, 1H), 7.32-7.26 (m, 1H), 2.80 (t, 2H), 2.64 (d, 3H), 2.50-2.45 (m, 2H), 1.65-1.56 (m, 2H), 1.55-1.45 (m, 2H), 1.33-1.26 (m, 15 4H).

Example 153

N~9~-(5-chloro-1,3-benzoxazol-2-yl)-N~1~-methyl-2-oxononanediamide

20 The desired product was prepared by substituting Example 124E and 5-chloro-2-aminobenzoxazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 366, 368 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 11.71 (s, 1H), 8.53-8.48 (m, 1H), 7.66-7.63 (m, 2H), 7.30 (dd, 1H), 2.80 (t, 2H), 2.64 (d, 3H), 2.50-2.46 (m, 2H), 1.63-1.45 (m, 4H), 1.32-1.26 (m, 4H).

25 Example 154

N~9~-(4-(4-chlorophenyl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-(4'-chlorophenyl)-2-aminothiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 408, 410 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 12.21 (s, 1H), 8.53-8.47 (m, 1H), 7.90 (d, 2H), 7.66 (s, 1H), 7.48 (d, 2H), 2.79 (t, 2H), 2.64 (d, 3H), 30 2.44 (t, 2H), 1.63-1.45 (m, 4H), 1.32-1.26 (m, 4H).

Example 155

N~9~-(4-(4-bromophenyl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide

35 The desired product was prepared by substituting Example 124E and 4-(4'-bromophenyl)-2-aminothiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 452, 454 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 12.22 (s,

1H), 8.53-8.47 (m, 1H), 7.84 (d, 2H), 7.67 (s, 1H), 7.62 (d, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.44 (t, 2H), 1.64-1.56 (m, 2H), 1.54-1.46 (m, 2H), 1.31-1.26 (m, 4H).

Example 156

N~9~-~(4-(3-bromophenyl)-1,3-thiazol-2-yl)-N~1~-~methyl-2-oxanonanediame

5 The desired product was prepared by substituting Example 124E and 4-(3'-bromophenyl)-2-aminothiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 452, 454 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 12.22 (s, 1H), 8.53-8.47 (m, 1H), 8.10-8.09 (m, 1H), 7.91-7.88 (m, 1H), 7.75 (s, 1H), 7.53-7.49 (m, 1H), 7.39 (t, 1H), 2.80 (t, 2H), 2.64 (d, 3H), 2.44 (t, 2H), 1.64-1.56 (m, 2H), 1.54-1.44 (m, 2H), 1.32-1.26 (m, 4H).

Example 157

N-(8-(methylamino)-7,8-dioxooctyl)-1,1'-biphenyl-3-carboxamide

15 The desired product was prepared by substituting Example 142F and 3-phenylbenzoic acid for aniline and Example 1B, respectively, in Example 1C. MS (ESI(+)) m/e 367 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.59-8.48 (m, 2H), 8.10 (t, 1H), 7.84-7.79 (m, 2H), 7.81-7.75 (m, 2H), 7.57-7.48 (m, 3H), 7.43-7.38 (m, 1H), 3.31-3.24 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.58-1.46 (m, 4H), 1.35-1.26 (m, 4H).

20

Example 158

3-(4-(1,1'-biphenyl-4-yloxy)butoxy)-N-methyl-2-oxopropanamide

Example 158A

4-(1,1'-biphenyl-4-yloxy)butan-1-ol

25 A solution of ethyl 4-((1,1'-biphenyl)-4-yloxy)butanoate (3.3 g, 11.6 mmol, prepared by substituting ethyl 4-bromobutanoate for ethyl 7-bromoheptanoate in Example 2A) in dichloromethane (100 mL) at -78 °C was treated with 1M DIBAL in dichloromethane (35 mL, 35 mmol), warmed to room temperature over 2 hours, quenched with saturated sodium 30 potassium tartrate, concentrated, and diluted with ethyl acetate. The aqueous phase was extracted twice with ethyl acetate and the combined organic phases were washed with water and brine, dried (Na₂SO₄), filtered, and concentrated to provide 2.8 g of the desired product. MS (ESI(+)) m/e 243 (M+H)⁺.

35

Example 158B

4-(4-(2,2-diethoxyethoxy)butoxy)-1,1'-biphenyl

A solution of Example 158A (2.8 g, 11.5 mmol) in DMF (40 mL) at 0 °C was treated with 60% NaH dispersion in oil (0.508 g, 12.7 mmol), warmed to room temperature, stirred for 2 hours, cooled to 0 °C, and treated dropwise with bromoacetaldehyde diethyl acetal (2.09 mL, 13.9 mmol). The mixture was heated to 90 °C for 18 hours, cooled to room temperature, and partitioned between water and ethyl acetate. The aqueous phase was extracted twice with ethyl acetate and the combined organic phases were washed with water and brine, dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 8:1 hexanes/ethyl acetate to provide 1.01g (24% yield) of the desired product. MS (ESI(+)) m/e 381 (M+Na)⁺.

10

Example 158C(4-(1,1'-biphenyl-4-yloxy)butoxy)acetaldehyde

A solution of Example 158B (1 g, 2.79 mmol) in 4:1 acetone/water (13 mL) was treated with conc. H₂SO₄ (9 drops), heated to reflux, stirred for 18 hours, cooled to room temperature, diluted with dichloromethane, washed with saturated NaHCO₃, dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 20:1 ethyl acetate/dichloromethane to provide the desired product as a mixture of aldehyde and hydrate. MS (DCI/NH₃) m/e 302 (M+NH₄)⁺.

20

Example 158Dmethyl 3-(4-(1,1'-biphenyl-4-yloxy)butoxy)-2-hydroxypropanoate

A suspension of Example 158C (100 mg, 0.35 mmol) in a mixture of water (1 mL) and THF (1 mL) at room temperature was treated with NaHSO₃ (57 mg) and KCN (34 mg), stirred for 18 hours, and concentrated under a stream of nitrogen. The resulting solid was collected by filtration and washed with cold water. The filter cake was dissolved in methanol (1 mL), cooled to 0 °C, treated with HBr gas for 1 hour, diluted with water (1 mL) and stirred for 30 minutes. The reaction was diluted with saturated NaHCO₃ and extracted three times with dichloromethane. The combined extracts were dried (Na₂SO₄), filtered, and concentrated. The concentrate was dissolved in methanol, stirred for 18 hours, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 1% methanol/dichloromethane to provide 80 mg (78% yield) of the desired product. MS (DCI/NH₃) m/e 362 (M+H)⁺.

Example 158E3-(4-(1,1'-biphenyl-4-yloxy)butoxy)-2-hydroxy-N-methylpropanamide

The desired product was prepared by substituting Example 158D for Example 101B in Example 101C.

Example 158F3-(4-(1,1'-biphenyl-4-yloxy)butoxy)-N-methyl-2-oxopropanamide

The desired product was prepared by substituting Example 158E for Example 86A in Example 86B. MS (DCI/NH₃) m/e 342 (M+H)⁺, 359 (M+NH₄)⁺; ¹H NMR (DMSO-d₆) δ 8.64-8.59 (m, 1H), 7.62-7.57 (m, 4H), 7.42 (t, 2H), 7.30 (t, 1H), 7.03-7.00 (m, 2H), 4.69 (s, 2H), 4.04 (t, 2H), 3.52 (t, 2H), 2.64 (d, 3H), 1.82-1.76 (m, 2H), 1.73-1.66 (m, 2H); Anal. Cald for C₂₀H₂₃NO₄·0.125H₂O: C, 69.90; H, 6.82; N, 4.08. Found: C, 69.91; H, 6.68; N, 3.90.

10

Example 159N-(8-(methylamino)-7,8-dioxooctyl)-2-phenyl-1,3-thiazole-4-carboxamide

The desired product was prepared by substituting Example 142F and 2-phenyl-4-thiazole carboxylic acid for 4-aminopyridine and Example 1B, respectively, in Example 6. MS (ESI(+)) m/e 374 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.53-8.47 (m, 2H), 8.26 (s, 1H), 8.08-8.04 (m, 2H), 7.56-7.51 (m, 3H), 3.31-3.25 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.59-1.46 (m, 4H), 1.33-1.28 (m, 4H).

Example 1605-(4-chlorophenyl)-N-(8-(methylamino)-7,8-dioxooctyl)-2-furamide

The desired product was prepared by substituting Example 142F and 5-(4-chlorophenyl)-2-furoic acid for 4-aminopyridine and Example 1B, respectively, in Example 6. MS (ESI(+)) m/e 391, 393 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.53-8.48 (m, 2H), 7.94 (d, 2H), 7.54 (d, 2H), 7.13 (s, 2H), 3.27-3.17 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.55-1.45 (m, 4H), 1.33-1.28 (m, 4H).

Example 1611-benzyl-N-(8-(methylamino)-7,8-dioxooctyl)-1H-indole-3-carboxamide

The desired product was prepared by substituting Example 142F and 1-benzyl-3-indole carboxylic acid for 4-aminopyridine and Example 1B, respectively, in Example 6. MS (ESI(+)) m/e 420 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.53-8.46 (br m, 1H), 8.16-8.13 (m, 1H), 8.09 (s, 1H), 7.88-7.85 (br m, 1H), 7.50 (d, 1H), 7.36-7.22 (m, 5H), 7.20-7.09 (m, 2H), 5.45 (s, 2H), 3.26-3.19 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.56-1.46 (m, 4H), 1.35-1.28 (m, 4H).

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Example 1625-(benzyloxy)-N-(8-(methylamino)-7,8-dioxooctyl)-1H-indole-2-carboxamide

The desired product was prepared by substituting Example 142F and 5-benzyloxy-2-indole carboxylic acid for 4-aminopyridine and Example 1B, respectively, in Example 6. MS (ESI(+)) m/e 420 (M-CH₃)⁺; ¹H NMR (DMSO-d₆) δ 11.37 (s, 1H), 8.53-8.48 (br m, 1H), 8.38-8.34 (br m, 1H), 7.49-7.46 (m, 2H), 7.42-7.30 (m, 4H), 7.16 (d, 1H), 6.99 (d, 1H), 6.90 (dd, 1H), 5.09 (s, 2H), 3.29-3.22 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.56-1.46 (m, 4H), 1.34-1.28 (m, 4H).

Example 163

N-(8-(methylamino)-7,8-dioxooctyl)benzamide

10 The desired product was prepared by substituting Example 142F and benzoic acid for 4-aminopyridine and Example 1B, respectively, in Example 6. MS (ESI(+)) m/e 291 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.54-8.48 (br m, 1H), 8.44-8.40 (br m, 1H), 7.84-7.81 (m, 2H), 7.51-7.41 (m, 3H), 3.27-3.20 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.54-1.66 (m, 4H), 1.32-1.27 (m, 4H).

15

Example 164

N~9~-(4-(4-cyanophenyl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide

10 The desired product was prepared by substituting Example 124E and 4-(4-cyanophenyl)-2-aminothiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 399 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.28 (s, 1H), 8.53-8.48 (br m, 1H), 8.07 (d, 2H), 7.90 (s, 1H), 7.89 (d, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.45 (t, 2H), 1.63-1.46 (m, 4H), 1.32-1.26 (m, 4H).

25

N~9~-(4-(2,3-dihydro-1-benzofuran-5-yl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide

10 The desired product was prepared by substituting Example 124E and 4-(2,3-dihydro-1-benzofuran-5-yl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 416 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.11 (s, 1H), 8.53-8.47 (br m, 1H), 7.75-7.74 (m, 1H), 7.65-7.62 (m, 1H), 7.36 (s, 1H), 6.79 (d, 1H), 4.56 (t, 2H), 3.21 (t, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.45 (t, 2H), 1.63-1.44 (m, 4H), 1.32-1.26 (m, 4H).

35

N~1~-methyl-2-oxo-N~9~-(4-(5,6,7,8-tetrahydronaphthalen-2-yl)-1,3-thiazol-2-yl)nonanediamide

The desired product was prepared by substituting Example 124E and 4-(5,6,7,8-tetrahydronaphthalen-2-yl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 428 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.17 (s, 1H), 8.54-8.48 (br m, 1H), 7.60-7.57 (m, 2H), 7.48 (s, 1H), 7.10-7.07 (m, 1H), 2.82-2.70 (m, 6H), 2.64 (d, 3H), 2.47-2.40 (m, 2H), 1.79-1.72 (m, 4H), 1.64-1.56 (m, 2H), 1.54-1.46 (m, 2H), 1.33-1.26 (m, 4H).

Example 167

10 N~9~-(4-(2,3-dihydro-1,4-benzodioxin-6-yl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-(2,3-dihydro-1,4-benzodioxin-6-yl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 432 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.13 (s, 1H), 8.54-8.47 (br m, 1H), 7.43 (s, 1H), 7.37-7.34 (m, 2H), 6.88 (d, 1H), 4.26 (app s, 4H), 2.80 (t, 2H), 2.64 (d, 3H), 2.43 (t, 2H), 1.63-1.45 (m, 4H), 1.32-1.26 (m, 4H).

Example 168

N~9~-(4-(2,4-dimethoxyphenyl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide

20 The desired product was prepared by substituting Example 124E and 4-(2,4-dimethoxyphenyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 434 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.07 (s, 1H), 8.53-8.47 (br m, 1H), 7.97 (d, 1H), 7.44 (s, 1H), 6.66-6.60 (m, 2H), 3.90 (s, 3H), 3.80 (s, 3H), 2.80 (t, 2H), 2.64 (d, 3H), 2.43 (t, 2H), 1.63-1.45 (m, 4H), 1.32-1.25 (m, 4H).

25 Example 169

N~9~-(4-(2,5-dimethoxyphenyl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-(2,5-dimethoxyphenyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 434 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.11 (s, 1H), 8.54-8.47 (br m, 1H), 7.66-7.63 (m, 2H), 7.05 (d, 1H), 6.90-6.85 (m, 1H), 3.86 (s, 3H), 3.74 (s, 3H), 2.80 (t, 2H), 2.64 (d, 3H), 2.47-2.42 (m, 2H), 1.62-1.46 (m, 4H), 1.31-1.26 (m, 4H).

Example 170

N~1~-methyl-2-oxo-N~9~-(4-(4-(trifluoromethyl)phenyl)-1,3-thiazol-2-yl)nonanediamide

35 The desired product was prepared by substituting Example 124E and 4-(trifluoromethyl)phenyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 442 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.28 (s,

1H), 8.53-8.48 (br m, 1H), 8.10 (d, 2H), 7.84-7.78 (m, 3H), 2.80 (t, 2H), 2.64 (d, 3H), 2.45 (t, 2H), 1.65-1.44 (m, 4H), 1.32-1.26 (m, 4H).

Example 171

5 N~9~-{(4-(1,1'-biphenyl-4-yl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide}

The desired product was prepared by substituting Example 124E and 4-(1,1'-biphenyl-4-yl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 450 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.22 (s, 1H), 8.53-8.48 (br m, 1H), 7.98 (d, 2H), 7.73-7.71 (m, 4H), 7.66 (s, 1H), 7.48 (t, 2H), 7.38-7.53 (m, 1H), 2.80 (t, 2H), 10 2.64 (d, 3H), 2.45 (t, 2H), 1.66-1.46 (m, 4H), 1.32-1.27 (m, 4H).

Example 172

N~1~-methyl-2-oxo-N~9~-{(4-(4-(trifluoromethoxy)phenyl)-1,3-thiazol-2-yl)nonanediamide}

The desired product was prepared by substituting Example 124E and 4-(4-(trifluoromethoxy)phenyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 458 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.24 (s, 1H), 8.54-8.48 (br m, 1H), 8.00 (d, 2H), 7.68 (s, 1H), 7.42 (d, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.47 (t, 2H), 1.62-1.56 (m, 2H), 1.53-1.45 (m, 2H), 1.31-1.26 (m, 4H).

20 Example 173

N~1~-methyl-2-oxo-N~9~-{(4-(3-(trifluoromethoxy)phenyl)-1,3-thiazol-2-yl)nonanediamide}

The desired product was prepared by substituting Example 124E and 4-(3-(trifluoromethoxy)phenyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 458 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.24 (s, 1H), 8.53-8.47 (br s, 1H), 7.95-7.91 (m, 1H), 7.86-7.83 (m, 1H), 7.79 (s, 1H), 7.51 (t, 1H), 25 7.33-7.29 (m, 1H), 2.80 (t, 2H), 2.64 (d, 3H), 2.45 (t, 2H), 1.62-1.57 (m, 2H), 1.53-1.46 (m, 2H), 1.31-1.27 (m, 4H).

Example 174

30 N~1~-methyl-2-oxo-N~9~-{(4-(3,4,5-trimethoxyphenyl)-1,3-thiazol-2-yl)nonanediamide}

The desired product was prepared by substituting Example 124E and 4-(3,4,5-trimethoxyphenyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 464 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.21 (s, 1H), 8.53-8.48 (br m, 1H), 7.61 (s, 1H), 7.91 (s, 2H), 3.83 (s, 6H), 3.68 (s, 3H), 2.80 (t, 2H), 2.64 (d, 3H), 2.44 (t, 2H), 1.62-1.46 (m, 4H), 1.31-1.26 (m, 4H).

Example 175

N~1~-methyl-2-oxo-N~9~- (4-(4-phenoxyphenyl)-1,3-thiazol-2-yl)nonanediamide

The desired product was prepared by substituting Example 124E and 4-(4-phenoxyphenyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 466 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.20 (s, 1H), 8.52-8.48 (br m, 1H), 7.89 (d, 2H), 7.52 (s, 1H), 7.44-7.39 (m, 2H), 7.20-7.14 (m, 1H), 7.08-7.03 (m, 4H), 2.80 (t, 2H), 2.64 (d, 3H), 2.44 (t, 2H), 1.64-1.56 (m, 2H), 1.54-1.46 (m, 2H), 1.31-1.26 (m, 4H).

Example 176N~9~- (4-(4-(benzyloxy)phenyl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-(benzyloxy)phenyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 480 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.15 (s, 1H), 8.53-8.47 (br m, 1H), 7.82 (d, 2H), 7.48-7.33 (m, 6H), 7.06 (d, 2H), 5.14 (s, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.43 (t, 2H), 1.63-1.54 (m, 2H), 1.53-1.46 (m, 2H), 1.31-1.27 (m, 4H).

Example 177N~1~-methyl-2-oxo-N~9~- (4-pyridin-3-yl-1,3-thiazol-2-yl)nonanediamide

The desired product was prepared by substituting Example 124E and 4-(pyridin-3-yl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 375 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.27 (s, 1H), 9.11 (d, 1H), 8.52 (dd, 1H), 8.52-8.48 (br m, 1H), 8.24-8.20 (m, 1H), 7.78 (s, 1H), 7.46 (dd, 1H), 2.80 (t, 2H), 2.64 (d, 3H), 2.45 (t, 2H), 1.65-1.56 (m, 2H), 1.54-1.45 (m, 2H), 1.32-1.27 (m, 4H).

Example 178N~1~-methyl-2-oxo-N~9~- (4-pyridin-4-yl-1,3-thiazol-2-yl)nonanediamide

The desired product was prepared by substituting Example 124E and 4-(pyridin-4-yl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 375 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.30 (s, 1H), 8.62 (d, 2H), 8.54-8.46 (br m, 1H), 7.96 (s, 1H), 7.83 (d, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.45 (t, 2H), 1.66-1.44 (m, 4H), 1.33-1.24 (m, 4H).

Example 179N~9~- (4-(3-cyanophenyl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-(3-cyanophenyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 399 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.25 (s, 1H), 8.53-8.48

(br m, 1H), 8.31 (br s, 1H), 8.21 (d, 1H), 7.84 (d, 1H), 7.78 (d, 1H), 7.65 (t, 1H), 2.80 (t, 2H), 2.64 (d, 3H), 2.45 (t, 2H), 1.65-1.45 (m, 4H), 1.32-1.26 (m, 4H).

Example 180

5 N~9~- (4-(4-ethoxyphenyl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-(4-ethoxyphenyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 418 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.15 (s, 1H), 8.52-8.48 (br m, 1H), 7.79 (d, 2H), 7.41 (s, 1H), 6.96 (d, 2H), 4.09-4.02 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.43 (t, 2H), 1.64-1.46 (m, 4H), 1.34 (t, 3H), 1.31-1.26 (m, 4H).

Example 181

N~1~-methyl-N~9~- (4-(2-naphthyl)-1,3-thiazol-2-yl)-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-(2-naphthyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 424 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.28 (s, 1H), 8.53-8.47 (br m, 1H), 8.42 (s, 1H), 8.06-8.03 (m, 1H), 7.97-7.90 (m, 3H), 7.75 (s, 1H), 7.75-7.49 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.46 (t, 2H), 1.67-1.46 (m, 4H), 1.33-1.28 (m, 4H).

20 Example 182

N~1~-methyl-N~9~- (4-(4-morpholin-4-ylphenyl)-1,3-thiazol-2-yl)-2-oxononanediamide

The desired product was prepared by substituting Example 124E and 4-(4-morpholin-4-ylphenyl)-2-amino-1,3-thiazole for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 459 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.13 (s, 1H), 8.52-8.47 (br m, 1H), 7.74 (d, 2H), 7.36 (s, 1H), 6.98 (d, 2H), 3.76-3.73 (m, 4H), 3.17-3.13 (m, 4H), 2.80 (t, 2H), 2.64 (d, 3H), 2.43 (t, 2H), 1.63-1.47 (m, 4H), 1.31-1.26 (m, 4H).

Example 183

N-methyl-2-oxo-8-((4-phenyl-1,3-thiazol-2-yl)thio)octanamide

30

Example 183A

6-((4-phenyl-1,3-thiazol-2-yl)sulfanyl)hexan-1-ol

The desired product was prepared by substituting 4-phenyl-1,3-thiazole-2-thiol and 6-bromohexanol for (1,1'-biphenyl)-4-ol and ethyl 7-bromoheptanoate, respectively, in Example 2A.

Example 183B

6-((4-phenyl-1,3-thiazol-2-yl)sulfanyl)hexanal

The desired product was prepared by substituting Example 183A for Example 142A in Example 142B.

5

Example 183Cmethyl (2Z)-2-((tert-butyl(dimethylsilyl)oxy)-8-((4-phenyl-1,3-thiazol-2-yl)sulfanyl)oct-2-enoate

The desired product was prepared by substituting Example 183B for Example 142B in Example 142C.

10

Example 183Dmethyl 2-oxo-8-((4-phenyl-1,3-thiazol-2-yl)sulfanyl)octanoate

The desired product was prepared by substituting Example 183C for Example 142C in Example 142D.

15

Example 183EN-methyl-2-oxo-8-((4-phenyl-1,3-thiazol-2-yl)thio)octanamide

The desired product was prepared by substituting Example 183D for Example 106A in Example 106B. mp: 107-108 °C; MS (ESI(-)) m/e 361 (M-H)⁻; ¹H NMR (DMSO-d₆) δ 8.50 (br s, 1H), 8.02 (s, 1H), 7.85-7.95 (m, 2H), 7.30-7.50 (m, 3H), 3.38 (t, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.20-1.80 (m, 8H); Anal. Calcd for: C₁₈H₂₂N₂O₂S₂: C, 59.64; H, 6.12; N, 7.73. Found: C, 59.59; H, 5.91; N, 7.44.

20

Example 1848-(1,3-benzothiazol-2-ylthio)-N-methyl-2-oxooctanamide

25

Example 184A6-(1,3-benzothiazol-2-ylsulfanyl)hexan-1-ol

The desired product was prepared by substituting 1,3-benzothiazole-2-thiol and 6-bromohexanol for (1,1'-biphenyl)-4-ol and ethyl 7-bromoheptanoate, respectively, in Example 2A.

30

Example 184B6-(1,3-benzothiazol-2-ylsulfanyl)hexanal

The desired product was prepared by substituting Example 184A for Example 142A in Example 142B.

35

Example 184C

methyl (2Z)-8-(1,3-benzothiazol-2-ylsulfanyl)-2-((tert-butyl(dimethyl)silyl)oxy)oct-2-enoate

The desired product was prepared by substituting Example 184B for Example 142B in Example 142C.

5

Example 184Dmethyl 8-(1,3-benzothiazol-2-ylsulfanyl)-2-oxooctanoate

The desired product was prepared by substituting Example 184C for Example 142C in Example 142D.

10

Example 184E8-(1,3-benzothiazol-2-ylthio)-N-methyl-2-oxooctanamide

The desired product was prepared by substituting Example 184D for Example 106A in Example 106B. mp: 83-84 °C; MS (ESI(+)) m/e 336.9 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.50 (s, 1H), 8.01 (dd, 1H), 7.85 (dd, 1H), 7.46 (dt, 1H), 7.47 (dt, 1H), 3.45 (m, 2H), 2.82 (t, 2H), 2.64 (d, 3H), 1.30-1.90 (m, 8H); Anal. Calcd for: C₁₆H₁₉NO₃S₂: C, 57.11; H, 5.99; N, 8.33. Found: C, 57.02; H, 5.89; N, 8.16.

15

Example 185N-(8-(methylamino)-7,8-dioxooctyl)-4-phenyl-1,3-thiazole-2-carboxamide

20

The desired product was prepared by substituting Example 142F and 4-phenyl-2-thiazolecarboxylic for aniline and Example 1B, respectively, in Example 1C. MS (ESI(+)) m/e 374 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.88-8.84 (br m, 1H), 8.52-8.47 (br m, 1H), 8.38 (s, 1H), 8.08 (d, 2H), 7.49 (t, 2H), 7.42-7.37 (m, 1H), 3.31-3.26 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.60-1.46 (m, 4H), 1.34-1.28 (m, 4H).

25

Example 186N~9~-{(1H-indol-5-yl)-N~1~-methyl-2-oxononanediamide}

30

The desired product was prepared by substituting Example 124E and 5-aminoindole for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+)) m/e 330 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 10.94 (br s, 1H), 9.61 (br s, 1H), 8.52-8.46 (br m, 1H), 7.85 (d, 1H), 7.29-7.26 (m, 2H), 7.17 (dd, 1H), 6.34 (t, 1H), 2.80 (t, 2H), 2.64 (d, 3H), 2.27 (t, 2H), 1.62-1.46 (m, 4H), 1.32-1.28 (m, 4H).

35

Example 187N~1~-methyl-2-oxo-N~9~-(3-phenyl-1,2,4-thiadiazol-5-yl)nonanediamide

The desired product was prepared by substituting Example 124E and 3-phenyl-1,2,4-thiadiazol-5-amine for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+))

m/e 375 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 13.04 (s, 1H), 8.53-8.47 (br m, 1H), 8.17-8.14 (m, 2H), 7.53-7.49 (m, 3H), 2.80 (t, 2H), 2.64 (d, 3H), 2.55 (t, 2H), 1.68-1.59 (m, 2H), 1.55-1.46 (m, 2H), 1.33-1.27 (m, 4H).

5

Example 188

N~1~-methyl-N~9~-{(1-methyl-5-phenyl-1H-pyrazol-3-yl)-2-oxanonanediamicide}

The desired product was prepared by substituting Example 124E and 1-methyl-5-phenyl-1H-pyrazol-3-amine for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+)) m/e 371 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.92 (br s, 1H), 8.53-8.47 (br m, 1H), 7.74 (d, 2H), 7.40-7.35 (m, 2H), 7.30-7.25 (m, 1H), 6.62 (s, 1H), 3.69 (s, 3H), 2.81 (t, 2H), 2.64 (d, 3H), 2.36 (t, 2H), 1.64-1.46 (m, 4H), 1.36-1.28 (m, 4H).

Example 189

N-methyl-2-oxo-8-((4-phenyl-1,3-thiazol-2-yl)sulfonyl)octanamide

The desired product was prepared by substituting Example 183 for Example 102D in Example 103. MS (ESI(-)) m/e 393 (M-H)⁻; ¹H NMR (DMSO-d₆) δ 8.66 (s, 1H), 8.50 (br, 1H), 7.95-8.05 (m, 2H), 7.40-7.60 (m, 2H), 3.63 (t, 2H), 2.78 (t, 2H), 2.64 (d, 3H). Anal. Calcd for: C₁₈H₂₂N₂O₄S₂: C, 54.80; H, 5.62; N, 7.10. Found: C, 54.23; H, 5.53; N, 6.93.

20

Example 190

N~1~-methyl-N~9~-{(4-(4-(2-morpholin-4-ylethoxy)phenyl)-1,3-thiazol-2-yl)-2-oxanonanediamicide}

The desired product was prepared by substituting Example 124E and 4-(4-(2-morpholin-4-ylethoxy)phenyl)-1,3-thiazol-2-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 503 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.14 (s, 1H), 8.53-8.48 (br m, 1H), 7.80 (d, 2H), 7.42 (s, 1H), 6.99 (d, 2H), 4.12 (t, 2H), 3.59-3.56 (m, 4H), 2.80 (t, 2H), 2.70 (t, 2H), 2.64 (d, 3H), 2.49-2.46 (m, 4H), 2.43 (t, 2H), 1.64-1.45 (m, 4H), 1.32-1.26 (m, 4H).

30

Example 191

N~1~-methyl-N~9~-{(4-(6-morpholin-4-ylpyridin-3-yl)-1,3-thiazol-2-yl)-2-oxanonanediamicide}

The desired product was prepared by substituting Example 124E and 4-(6-morpholin-4-ylpyridin-3-yl)-1,3-thiazol-2-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 460 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.17 (s, 1H), 8.66 (d, 1H), 8.53-8.48 (br m, 1H), 8.01 (dd, 1H), 7.43 (s, 1H), 6.89 (d, 1H), 3.79-3.62 (m, 4H), 3.50-3.47 (m, 4H), 2.80 (t, 2H), 2.64 (d, 3H), 2.43 (t, 2H), 1.62-1.44 (m, 4H), 1.31-1.25 (m, 4H).

Example 192N~9~-(4-(4-(2-(dimethylamino)ethoxy)phenyl)-1,3-thiazol-2-yl)-N~1~-methyl-2-oxononanediamide

5 The desired product was prepared by substituting Example 124E and 4-(4-(2-(dimethylamino)ethoxy)phenyl)-1,3-thiazol-2-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 461 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.14 (s, 1H), 8.53-8.47 (br m, 1H), 7.80 (d, 2H), 7.41 (s, 1H), 6.98 (d, 2H), 4.07 (t, 2H), 2.77 (t, 2H), 2.65-2.61 (m, 5H), 2.43 (t, 2H), 2.22 (s, 6H), 1.62-1.47 (m, 4H), 1.31-1.26 (m, 4H).

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Example 193N~1~-methyl-N~9~-(4-(4-(4-methylpiperazin-1-yl)phenyl)-1,3-thiazol-2-yl)-2-oxononanediamide

15 The desired product was prepared by substituting Example 124E and 4-(4-(4-methylpiperazin-1-yl)phenyl)-1,3-thiazol-2-amine for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 472 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.12 (s, 1H), 8.53-8.47 (br m, 1H), 7.72 (d, 2H), 7.33 (s, 1H), 6.96 (d, 2H), 3.19-3.16 (m, 4H), 2.79 (t, 2H), 2.64 (d, 3H), 2.47-2.40 (m, 6H), 2.23 (s, 3H), 1.62-1.47 (m, 4H), 1.31-1.27 (m, 4H).

20

Example 194N-methyl-8-(((2-naphthylamino)carbonyl)amino)-2-oxooctanamide

A solution of 2-naphthylisocyanate (47 mg, 0.27 mmol) in dichloromethane (6 mL) at room temperature was treated with Example 142F (60 mg, 0.27 mmol) and triethylamine, (0.042 mL), stirred at room temperature for 2 hours, and partitioned between 25 dichloromethane and 1N HCl. The organic phase was washed with brine, diluted with methanol, dried (Na₂SO₄), filtered, and concentrated to provide 34 mg of the desired product. MS (ESI(+)) m/e 356 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.60 (d, 1H), 8.01 (s, 1H), 7.79-7.68 (m, 3H), 7.36-7.46 (m, 2H), 7.27-7.34 (m, 1H), 6.21 (t, 1H), 3.10 (dt, 2H), 2.81 (t, 2H), 2.64 (d, 3H), 1.38-1.57 (m, 4H), 1.25-1.57 (m, 4H); Anal. Calcd for C₂₀H₂₅N₃O₃: C, 67.58; H, 7.09; N, 11.82. Found: C, 67.34; H, 6.91; N, 11.70.

30

Example 1958-(5-(4-methoxyphenyl)-1,3-oxazol-2-yl)-N-methyl-2-oxooctanamide

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Example 195Amethyl 7-((2-(4-methoxyphenyl)-2-oxoethyl)amino)-7-oxoheptanoate

The desired product was prepared by substituting 7-methoxy-7-oxoheptanoic acid and 2-amino-4'-methoxyacetophenone for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+)) m/e 322 (M+H)⁺.

5

Example 195Bmethyl 6-(5-(4-methoxyphenyl)-1,3-oxazol-2-yl)hexanoate

A solution of Example 195A (1.1 g, 3.43 mmol) in CHCl₃ (30 mL) was treated with P₂O₅ (3.89 g, 13.7 mmol), heated to reflux for 18 hours, cooled to room temperature, and partitioned between water and dichloromethane. The aqueous phase was extracted twice with dichloromethane and the combined organic phases were dried (MgSO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography with 1:1 hexanes/ethyl acetate to provide 0.71g (68% yield) of the desired product. MS (ESI(+)) m/e 304 (M+H)⁺.

15

Example 195C6-(5-(4-methoxyphenyl)-1,3-oxazol-2-yl)hexan-1-ol

The desired product was prepared by substituting Example 195B for ethyl 4-(1,1'-biphenyl)-4-yloxy)butanoate in Example 158A.

20

Example 195D6-(5-(4-methoxyphenyl)-1,3-oxazol-2-yl)hexanal

The desired product was prepared by substituting Example 195C for Example 86A in Example 86B. MS (CI) m/e 274 (M+H)⁺.

25

Example 195Emethyl 8-(5-(4-methoxyphenyl)-1,3-oxazol-2-yl)-2-oxooctanoate

The desired product was prepared by substituting Example 195D for Example 142B in Examples 142C and 142D. MS (ESI(+)) m/e 346 (M+H)⁺.

30

Example 195F8-(5-(4-methoxyphenyl)-1,3-oxazol-2-yl)-N-methyl-2-oxooctanamide

The desired product was prepared by substituting Example 195E for Example 106A in Example 106B. MS (ESI(+)) m/e 345(M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.50 (br s, 1H), 7.59(d, 2H), 7.36 (s, 1H), 7.02(d, 2H), 3.79 (s, 3H), 2.79 (t, 2H), 2.77 (t, 2H), 2.64 (d, 3H), 1.71 (m, 2H), 1.50 (m, 1H), 1.32 (m, 4H); Anal. Calcd. for C₁₉H₂₄N₂O₄: C, 66.26; H, 7.02; N, 8.13. Found: C, 65.91; H, 6.92; N, 7.86.

Example 196N-methyl-2-oxo-8-(2-phenyl-1,3-thiazol-4-yl)octanamideExample 196A

5

methyl 8-bromo-7-oxoheptanoate

A solution of 7-methoxy-7-oxoheptanoic acid (4.6 g, 26.6 mmol) in dichloromethane at room temperature (200 mL) was treated with oxalyl chloride (2.55 mL) and 1 drop of DMF, stirred for 1 hour, concentrated, and dissolved in diethyl ether (2 mL) to provide solution A. A mixture of diethyl ether (150 mL) and 40% aqueous KOH (45 mL) at 0 °C was 10 treated portionwise with 1-methyl-3-nitro-1-nitrosoquanidine (15 g), and stirred for 10 minutes. The organic phase was dried over KOH, filtered, cooled to 0 °C, treated with solution A, stirred at 0 °C for 1.5 hours, treated with conc. HBr (33 mL), warmed to room temperature, and stirred for 30 minutes. The reaction was partitioned between water and ethyl acetate and the organic phase was washed with saturated NaHCO₃, dried (Na₂SO₄), 15 filtered and concentrated to provide 5.69g (85% yield) of the desired product. MS (DCI) m/e 268, 270 (M+NH₄)⁺.

Example 196Bmethyl 6-(2-phenyl-1,3-thiazol-4-yl)hexanoate

20 A mixture of thiobenzamide (0.546g) and Example 196A 91 g, 3.98 mmol) in methanol (20 mL) at room temperature was stirred for 18 hours, concentrated, then partitioned between aq. NaHCO₃ and ethyl acetate. The organic phase was dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel with 10% ethyl acetate/hexanes to provide 0.9g (78%) of the desired product. MS 25 (ESI(+)) m/e 290 (M+H)⁺.

Example 196C6-(2-phenyl-1,3-thiazol-4-yl)hexanal

A -78 °C solution of Example 196B (0.7 g, 2.42 mmol) in dichloromethane (15 mL) 30 was treated with 1M DIBAL in toluene (6.05 mL), stirred for 1 hour, treated with additional DIBAL (3 mL), stirred for 30 minutes, and quenched with methanol (0.7 mL) and Rochelle's salt. The reaction was warmed to room temperature and extracted three times with ethyl acetate. The combined extracts were washed with brine, dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel 35 with 20-30% ethyl acetate/hexanes to provide 0.47g (75%) of the desired product. MS (ESI(+)) m/e 260 (M+H)⁺.

Example 196Dmethyl (2Z)-2-((tert-butyl(dimethyl)silyloxy)-8-(2-phenyl-1,3-thiazol-4-yl)oct-2-enoate

The desired product was prepared by substituting Example 196C for Example 142B in Example 142C.

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Example 196Emethyl 2-oxo-8-(2-phenyl-1,3-thiazol-4-yl)octanoate

The desired product was prepared by substituting Example 196D for Example 142C in Example 142D.

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Example 196FN-methyl-2-oxo-8-(2-phenyl-1,3-thiazol-4-yl)octanamide

The desired product was prepared by substituting Example 196E for Example 106A in 106B. mp: 70-72 °C; MS (ESI(+)) m/e 331 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 1.32-1.34 (m, 4H), 1.46-1.55 (m, 2H), 1.64-1.74 (m, 2H), 2.63-2.65 (d, 3H), 2.72-2.82 (m, 4H), 7.33 (s, 1H), 7.46-7.51 (m, 3H), 7.89-7.93 (m, 2H), 8.46-8.54 (br s, 1H); Anal. Calcd for: C₁₈H₂₂N₂O₂S C, 65.43; H, 6.71; N, 8.48. Found: C, 65.27; H, 6.64; N, 8.26.

Example 1978-((1,1'-biphenyl-4-ylsulfonyl)amino)-N-methyl-2-oxooctanamide

The desired product was prepared by substituting Example 142F and 4-phenylbenzenesulfonyl chloride for Example 143A and 4-chlorophenylsulfonyl chloride in Example 145. MS (ESI(-)) m/e 401 (M-H)⁻; ¹H NMR (DMSO-d₆) δ 8.52-8.47 (br m, 1H), 7.90-7.83 (m, 4H), 7.73 (d, 2H), 7.62-7.58 (br m, 1H), 7.54-7.49 (m, 2H), 7.46-7.41 (m, 1H), 2.79-2.73 (m, 4H), 2.64 (d, 3H), 1.47-1.31 (m, 4H), 1.31-1.25 (m, 4H).

Example 1987-(((1E)-1,1'-biphenyl-4-ylmethyldene)amino)oxy)-N-methyl-2-oxoheptanamide

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Example 198A1,1'-biphenyl-4-carbaldehyde oxime

A mixture of 4-phenylbenzaldehyde (3.64 g, 20 mmol), NH₂OH·HCl (2.72 g, 40 mmol) and pyridine (1 mL) in THF (20 mL) and ethanol (20 mL) was heated to reflux for 2 hours, cooled to room temperature, and concentrated. The residue was suspended in water and filtered. The filter cake was washed with water, and dried to provide 3.9g of the desired product. MS (ESI(+)) m/e 198 (M+H)⁺.

Example 198B1,1'-biphenyl-4-carbaldehyde O-(5-hydroxypentyl)oxime

The desired product was prepared by substituting Example 198A and 5-(t-butyldimethylsilyloxy)-pentyl bromide (1,1'-biphenyl)-4-ol and 6-(t-butyldimethylsilyloxy)-hexyl bromide, respectively, in Example 91A.

Example 198C1,1'-biphenyl-4-carbaldehyde O-(5-oxopentyl)oxime

The desired product was prepared by substituting Example 198B for Example 142A
10 in Example 142B.

Example 198Dmethyl (2Z)-7-(((1E)-1,1'-biphenyl-4-ylmethylene)amino)oxy)-2-((tert-butyl(dimethylsilyl)oxy)hept-2-enoate

15 The desired product was prepared by substituting Example 198C for Example 142B in Example 142C.

Example 198Emethyl 7-(((1E)-1,1'-biphenyl-4-ylmethylene)amino)oxy)-2-oxoheptanoate

20 The desired product was prepared by substituting Example 198D for Example 142C in Example 142D.

Example 198F7-(((1E)-1,1'-biphenyl-4-ylmethylidene)amino)oxy)-N-methyl-2-oxoheptanamide

25 The desired product was prepared by substituting Example 198E for Example 106A in Example 106B. mp: 86-87 °C. MS (ESI(+)) m/e 353 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.52 (br s, 1H), 8.29 (s, 1H), 7.30-7.80 (m, 9H), 4.12 (t, 2H), 2.84 (t, 2H), 1.30-1.80 (m, 6H).

Example 199N-methyl-2-oxo-8-(2-phenyl-1,3-oxazol-5-yl)octanamideExample 199Amethyl 8-(bis(tert-butoxycarbonyl)amino)-7-oxooctanoate

A mixture of Example 196A (4.69 g, 18.7 mmol) and (Boc)₂NK (5.24 g, prepared
35 according to the procedure described in *J.Chem. Soc. Perkin Trans. 1983*, 2983) in DMF (50 mL) was heated to 90 °C for 1 hour, then partitioned between water and ethyl acetate. The aqueous phase was extracted with ethyl acetate twice and the combined extracts were washed with water, brine, dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified

by flash column chromatography on silica gel with 20% ethyl acetate/hexanes to provide 6.57 g (91%) of the desired product. MS (ESI(+)) m/e 410 (M+Na)⁺.

Example 199B

5

methyl 8-amino-7-oxo octanoate

A mixture of Example 199A (6.57 g) and 4N HCl in dioxane (20 mL) was stirred at room temperature for 1 hour, then diluted with diethyl ether. The precipitate was collected by filtration and the filter cake was dried to provide 3.22g (80%) of the desired product. MS (ESI(+)) m/e 188 (M+H)⁺.

10

Example 199C

methyl 6-(2-phenyl-1,3-oxazol-5-yl)hexanoate

The desired product was prepared by substituting Example 199B and benzoic acid for aniline and Example 1B, respectively, in Example 1C.

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Example 199D

methyl 6-(2-phenyl-1,3-oxazol-5-yl)hexanoate

The desired product was prepared by substituting Example 199C for Example 195A in Example 195B.

20

Example 199E

N-methyl-2-oxo-8-(2-phenyl-1,3-oxazol-5-yl)octanamide

The desired product was prepared by substituting Example 199D for Example 196B in Examples 196C and 196D. mp, 78-80 °C; MS (ESI(+)) m/e 315 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 1.26-1.40 (m, 4H), 1.46-1.55 (m, 2H), 1.60-1.70 (m, 2H), 2.63-2.65 (d, 3H), 2.70-2.75 (t, 2H), 2.78-2.82 (t, 2H), 7.01 (s, 1H), 7.45-7.54 (m, 3H), 7.91-7.94 (m, 2H), 8.5 (br s, 1H); Anal. Calcd for: C₁₈H₂₂N₂O₃: C, 68.77; H, 7.05; N, 8.91. Found: C, 68.49; H, 7.01; N, 8.67.

30

Examples 200-215 were prepared in parallel using a Quest Apparatus. Each vessel was charged with 40 mg of Example 142F, 1.5 equivalents of the appropriate isocyanate (the isocyanates used are listed in each example), 0.043 mL of triethylamine and 3 mL of DMF. The reactions were mixed for 2 hours, then treated with PS-Trisamine resin (0.27 mmol), and mixed for an additional 2 hours. The reaction vessels were filtered and rinsed into scintillation vials and concentrated on a high speed vacuum centrifuge. The residues were then purified by preparative HPLC with a gradient system of 0 to 95% acetonitrile in water (containing 0.1% TFA) over 10 minutes to provide the desired products.

Example 2008-((anilinocarbonyl)amino)-N-methyl-2-oxooctanamide

5 Isocyanate: isocyanato benzene. MS (ESI(+)) m/e 306 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.50 (s, 1H), 8.33 (s, 1H), 7.33-7.40 (m, 2H), 7.17-7.23 (m, 2H), 6.83-6.90 (m, 1H), 6.08 (t, 1H), 3.05 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.23-1.56 (m, 8H); Anal. Calcd for C₁₆H₂₃N₃O₃·0.25 H₂O: C, 62.02; H, 7.64; N, 13.56. Found: C, 62.12; H, 7.24; N, 13.58.

Example 20110 N-methyl-8-(((2-methylphenyl)amino)carbonyl)amino)-2-oxooctanamide

15 Isocyanate: 1-isocyanato-2-methylbenzene. MS (ESI(+)) m/e 320 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.51 (s, 1H), 7.80 (d, 1H), 7.54 (s, 1H), 7.02-7.12 (m, 2H), 6.81-6.88 (m, 1H), 6.50 (t, 1H), 3.07 (dt, 2H), 2.81 (t, 2H), 2.65 (d, 3H), 2.16 (s, 3H), 1.25-1.57 (m, 8H); Anal. Calcd for C₁₇H₂₅N₃O₃·0.25 H₂O: C, 63.04; H, 7.93; N, 12.97. Found: C, 63.37; H, 7.72; N, 12.97.

Example 202N-methyl-8-(((3-methylphenyl)amino)carbonyl)amino)-2-oxooctanamide

20 Isocyanate: 1-isocyanato-3-methylbenzene. MS (ESI(+)) m/e 320 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.50 (s, 1H), 8.26 (s, 1H), 7.21 (s, 1H), 7.03-7.18 (m, 2H), 6.69 (d, 1H), 6.07 (t, 1H), 3.05 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.13 (s, 3H), 1.22-1.55 (m, 8H); Anal. Calcd for C₁₇H₂₅N₃O₃: C, 63.93; H, 7.89; N, 13.16. Found: C, 63.72; H, 7.69; N, 13.05.

25

Example 203N-methyl-8-(((4-methylphenyl)amino)carbonyl)amino)-2-oxooctanamide

30 Isocyanate: 1-isocyanato-3-methylbenzene. MS (ESI(+)) m/e 320 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.50 (s, 1H), 8.22 (s, 1H), 7.21-7.28 (m, 2H), 7.00 (d, 2H), 6.02 (t, 1H), 3.04 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 2.20 (s, 3H), 1.22-1.56 (m, 8H); Anal. Calcd for C₁₇H₂₅N₃O₃: C, 63.93; H, 7.89; N, 13.16. Found: C, 63.55; H, 7.66; N, 12.95.

Example 2048-(((2-methoxyphenyl)amino)carbonyl)amino)-N-methyl-2-oxooctanamide

35 Isocyanate: 1-isocyanato-2-methoxybenzene. MS (ESI(+)) m/e 336 (M+H)⁺;

¹H NMR (300 MHz, DMSO-d₆) δ 8.50 (s, 1H), 8.04-8.10 (m, 1H), 7.84 (s, 1H), 6.91-6.97 (m, 1H), 6.78-6.89 (m, 3H), 3.82 (s, 3H), 3.05 (dt, 2H), 2.80 (t, 2H), 2.65 (d, 3H), 1.22-1.56 (m, 8H).

Anal. Calcd for C₁₇H₂₅N₃O₄: C, 60.88; H, 7.51; N, 12.53. Found: C, 60.49; H, 7.31; N,

5 12.39.

Example 205

8-(((3-methoxyphenyl)amino)carbonyl)amino)-N-methyl-2-oxooctanamide

Isocyanate: 1-isocyanato-3-methoxybenzene. MS (ESI(+)) m/e 336 (M+H)⁺;

10 ¹H NMR (300 MHz, DMSO-d₆) δ 8.52 (m, 1H), 8.38 (s, 1H), 7.06-7.15 (m, 2H), 6.83 (m, 1H), 6.45 (dd, 1H), 6.09 (t, 1H), 3.69 (s, 3H), 3.05 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.22-1.56 (m, 8H); Anal. Calcd for C₁₇H₂₅N₃O₄: C, 60.88; H, 7.51; N, 12.53. Found: C, 60.62; H, 7.32; N, 12.42.

15

Example 206

8-(((4-methoxyphenyl)amino)carbonyl)amino)-N-methyl-2-oxooctanamide

Isocyanate: 1-isocyanato-4-methoxybenzene. MS (ESI(+)) m/e 336 (M+H)⁺; ¹H

NMR (300 MHz, DMSO-d₆) δ 8.52 (m, 1H), 8.16 (s, 1H), 7.27 (m, 2H), 6.79 (m, 2H), 5.99 (t, 1H), 3.68 (s, 3H), 3.03 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H, 5.1 Hz), 1.22-1.55 (m, 8H);

20 Anal. Calcd for C₁₇H₂₅N₃O₄: C, 60.88; H, 7.51; N, 12.53. Found: C, 60.75; H, 7.21; N, 12.45.

Example 207

8-(((4-chlorophenyl)amino)carbonyl)amino)-N-methyl-2-oxooctanamide

Isocyanate: 1-isocyanato-4-chlorobenzene. MS (ESI(+)) m/e 340.7 (M+H)⁺; ¹H NMR

25 (300 MHz, DMSO-d₆) δ 8.48-8.52 (m, 2H), 7.37-7.43 (m, 2H), 7.21-7.27 (m, 2H), 6.14 (t, 1H), 3.05 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.23-1.54 (m, 8H); Anal. Calcd for C₁₆H₂₂N₃O₃Cl: C, 56.55; H, 6.53; N, 12.37. Found: C, 56.50; H, 6.40; N, 12.33.

30

Example 208

8-(((3-bromophenyl)amino)carbonyl)amino)-N-methyl-2-oxooctanamide

Isocyanate: 1-isocyanato-3-bromobenzene. MS (ESI(+)) m/e 383.9, 385.9 (M+H)⁺;

¹H NMR (300 MHz, DMSO-d₆) δ 8.60 (s, 1H), 8.53 (m, 1H), 7.81 (t, 1H), 7.12-7.23 (m, 2H), 7.01-7.07 (m, 1H), 6.20 (t, 1H), 3.05 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.23-1.55 (m, 8H).

35

Example 209

8-(((4-(dimethylamino)phenyl)amino)carbonyl)amino)-N-methyl-2-oxooctanamide

The desired product was prepared as the trifluoroacetate salt using 4-isocyanato-N,N-dimethylaniline. MS (ESI(+)) m/e 349.2 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.50 (m, 1H), 8.40 (m, 1H), 7.32-7.41 (m, 2H), 7.02-7.21 (m, 2H), 6.10 (m, 1H), 2.93-3.10 (m, 8H), 2.80 (t, 2H), 2.44 (d, 3H), 1.22-1.55 (m, 8H); Anal. Calcd for
 5 C₁₈H₂₈N₄O₃·CF₃CO₂H·H₂O: C, 49.99; H, 6.50; N, 11.66. Found: C, 49.96; H, 6.23; N, 11.61.

Example 210

N-methyl-2-oxo-8-(((3-(trifluoromethyl)phenyl)amino)carbonyl)amino)octanamide

Isocyanate: 1-isocyanato-3-trifluoromethylbenzene. MS (ESI(+)) m/e 374 (M+H)⁺;
 10 ¹H NMR (300 MHz, DMSO-d₆) δ 8.78 (s, 1H), 8.53 (m, 1H), 7.97 (s, 1H), 7.39-7.51 (m, 2H), 7.21 (d, 1H), 6.26 (t, 1H), 3.07 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.23-1.55 (m, 8H).

Example 211

N-methyl-2-oxo-8-(((3-phenoxyphenyl)amino)carbonyl)amino)octanamide

Isocyanate: 1-isocyanato-3-phenoxybenzene. MS (ESI(+)) m/e 398 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.53 (m, 1H), 8.39 (s, 1H), 7.30-7.42 (m, 4H), 7.02-7.10 (m, 1H), 6.89-6.95 (m, 4H), 6.08 (t, 1H), 3.06 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.24-1.55 (m, 8H); Anal. Calcd for C₂₂H₂₇N₃O₄: C, 66.48; H, 6.85; N, 10.57. Found: C, 66.33; H, 6.80; N, 10.45.

20

Example 212

N-methyl-2-oxo-8-(((4-phenoxyphenyl)amino)carbonyl)amino)octanamide

Isocyanate: 1-isocyanato-4-phenoxybenzene. MS (ESI(+)) m/e 398 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.51 (m, 1H), 8.38 (s, 1H), 7.30-7.42 (m, 4H), 7.02-7.09 (m, 1H), 6.88-6.95 (m, 4H), 6.08 (t, 1H), 3.06 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.25-1.56 (m, 8H); Anal. Calcd for C₂₂H₂₇N₃O₄: C, 66.48; H, 6.85; N, 10.57. Found: C, 66.35; H, 6.81; N, 10.49.

Example 213

8-(((1,1'-biphenyl-2-ylamino)carbonyl)amino)-N-methyl-2-oxooctanamide

Isocyanate: 2-isocyanato-1,1'-biphenyl. MS (ESI(+)) m/e 382 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.52 (m, 1H), 7.87 (d, 1H), 7.01-7.52 (m, 9H), 6.54 (t, 1H), 3.00 (dt, 2H), 2.79 (t, 2H), 2.64 (d, 3H), 1.20-1.53 (m, 8H).

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Example 214

8-(((3,5-dimethoxyphenyl)amino)carbonyl)amino)-N-methyl-2-oxooctanamide

Isocyanate: 1-isocyanato-3,5-dimethoxybenzene. MS (ESI(+)) m/e 366.6 (M+H)⁺; ¹H NMR (500 MHz, DMSO-d₆) δ 8.48 (m, 1H), 8.35 (s, 1H), 6.61 (m, 2H), 6.03-6.08 (m, 2H), 3.67 (s, 6H), 3.04 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.45-1.58 (m, 2H), 1.34-1.45 (m, 2H), 1.23-1.32 (m, 4H); Anal. Calcd for C₁₈H₂₇N₃O₅: C, 59.16; H, 7.45; N, 11.50. Found: C, 5 59.18; H, 7.06; N, 11.38.

Example 215

8-(((1,3-benzodioxol-5-ylamino)carbonyl)amino)-N-methyl-2-oxooctanamide

Isocyanate: 5-isocyanato-1,3-benzodioxole. MS (ESI(+)) m/e 350 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.50 (m, 1H), 8.24 (s, 1H), 7.16 (m, 1H), 6.73-6.77 (m, 1H), 6.62-6.66 (m, 1H), 6.00 (m, 1H), 5.91 (s, 2H), 3.03 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.23-1.54 (m, 8H); Anal. Calcd for C₁₇H₂₃N₃O₅: C, 58.44; H, 6.63; N, 12.03. Found: C, 58.52; H, 6.28; N, 12.00.

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Example 216

N~7~-methyl-7-oxo-N~1~,N~1~-diphenylheptane-1,1,7-tricarboxamide

Example 216A

di-tert-butyl 2-(5-((tert-butyl(dimethyl)silyl)oxy)pentyl)malonate

20 A suspension of 95% NaH oil dispersion (380 mg, 15 mmol) in THF (50 mL) at 0 °C was treated dropwise with di-tert-butyl malonate (2.65 mL, 11.8 mmol), warmed to room temperature over 30 minutes, treated with 5-(t-butyldimethylsilyloxy)pentyl bromide (3.30 g, 11.7 mmol), heated to reflux for 18 hours, and partitioned between water and diethyl ether. The organic phase was washed with brine, dried (Na₂SO₄), filtered, concentrated and purified 25 by flash column chromatography on silica gel with 2 to 3% ethyl acetate/hexanes to provide 1.66g (34%) of the desired product. MS (ESI(-)) m/e 415 (M-H)⁻.

Example 216B

di-tert-butyl 2-(5-hydroxypentyl)malonate

30 A solution of Example 216A (1.66 g, 4.2 mmol) in THF (1 mL) was treated with 1M TBAF in THF (8.5 mL, 8.5 mmol), stirred for 3 hours, and partitioned between water and diethyl ether. The organic phase was washed with brine, dried (MgSO₄), filtered, concentrated, and purified by flash column chromatography on silica gel with 20% ethyl acetate/hexanes to provide 0.87 g (69%) of the desired product. MS (ESI(-)) m/e 301 (M-H)⁻.

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Example 216C

1,1-di-tert-butyl 7-methyl 7-oxoheptane-1,1,7-tricarboxylate

The desired product was prepared by substituting Example 216B for Example 142A in Examples 142B, 142C, and 142D.

Example 216D

5 di-tert-butyl 2-(7-(methylamino)-6,7-dioxoheptyl)malonate

The desired product was prepared by substituting Example 216C for Example 106A in Example 106B. MS (ESI(-)) m/e 370 (M-H)⁻.

Example 216E

10 2-(7-(methylamino)-6,7-dioxoheptyl)malonic acid

A solution of Example 216C (0.516 g, 1.4 mmol) in HCOOH (16 mL) at room temperature was stirred for 8 hours and concentrated. The reaction was concentrated under a stream of nitrogen to provide 0.364g of the desired product. MS (ESI(-)) m/e 258 (M-H)⁻.

15 Example 216F

N~7~-methyl-7-oxo-N~1~,N~1~-diphenylheptane-1,1,7-tricarboxamide

The desired product was prepared by substituting Example 216E for Example 1B in Example 1C. MS (ESI(+)) m/e 410 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.91 (s, 2H), 8.52-8.46 (br m, 1H), 7.60 (d, 4H), 7.31 (m, 4H), 7.05 (t, 2H), 3.49-3.45 (m, 1H), 2.79 (t, 2H), 2.63 (d, 3H), 1.94-1.85 (br m, 2H), 1.54-1.46 (m, 2H), 1.33-1.28 (m, 4H).

Example 217

8-(2-(4-bromophenyl)-1,3-oxazol-5-yl)-N-methyl-2-oxooctanamide

The desired product was prepared by substituting 4-bromobenzoic acid for benzoic acid in Example 199. mp: 90-92 °C; MS (ESI(+)) m/e 393, 395 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 1.27-1.41 (m, 4H), 1.43-1.57 (m, 2H), 1.57-1.70 (m, 2H), 2.63-2.65(d, 2H), 2.70-2.74 (t, 2H), 2.77-2.82 (t, 2H), 7.04 (1H), 7.70-7.73 (d, 2H), 7.84-7.87 (d, 2H), 8.50 (s, 1H); Anal. Calcd for: C₁₈H₂₁BrN₂O₃: C, 54.97; H, 5.38; N, 7.12. Found: C, 54.78; H, 5.46; N, 6.91.

30 Example 218

8-(2-(4-chlorophenyl)-1,3-thiazol-4-yl)-N-methyl-2-oxooctanamide

The desired product was prepared by substituting 4-chlorothiobenzamide for thiobenzamide in Example 196. mp: 77-81 °C; MS (ESI(+)) m/e 365, 367 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 1.23-1.40 (m, 4H), 1.45-1.55 (m, 2H), 1.63-1.73 (m, 2H), 2.63-2.65 (d, 3H), 2.72-2.82 (m, 4H), 7.37 (s, 1H), 7.53-7.56 (d, 2H), 7.91-7.94 (d, 2H), 8.49 (br s, 1H); Anal. Calcd for: C₁₈H₂₁ClN₂O₂S: C, 59.25; H, 5.80; N, 7.68. Found: C, 59.29; H, 5.86; N, 7.45.

Example 219methyl 9-(1,1'-biphenyl-3-ylamino)-3-hydroxy-2,9-dioxononanoateExample 219A5 N-1,1'-biphenyl-3-yl-7-hydroxyheptanamide

The desired product was prepared by substituting 1,1'-biphenyl-3-amine and 8-hydroxyoctanoic acid for anilin and Example 1B, respectively in Example 1C.

Example 219B10 N-1,1'-biphenyl-3-yl-7-oxoheptanamide

The desired product was prepared by substituting Example 219A for Example 142A in Example 142B.

Example 219C15 methyl 9-(1,1'-biphenyl-3-ylamino)-3-hydroxy-2,9-dioxononanoate

The desired product was prepared by substituting Example 219B for Example 141A in Examples 141B, 141C, and 141D. MS (DCI) m/e 384 (M+H)⁺, 401 (M+NH₄)⁺; ¹H NMR (DMSO-d₆) δ 9.94 (s, 1H), 7.92 (br s, 1H), 7.61-7.55 (m, 3H), 7.47 (d, 2H), 7.40-7.29 (m, 3H), 6.82 (d, 1H), 4.20 (br d, 1H), 3.69 (s, 3H), 2.30 (t, 2H), 1.61-1.51 (m, 2H), 1.41-1.16 (m, 6H); Anal Calcd for C₂₂H₂₅NO₅·0.25H₂O: C, 68.11; H, 6.63; N, 3.61. Found: C, 68.26; H, 6.55; N, 3.47.

Example 220methyl 9-((4-(4-methoxyphenyl)-1,3-thiazol-2-yl)amino)-2,9-dioxononanoate

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Example 220A6-(1,3-dioxolan-2-yl)hexanoic acid

The desired product was by substituting methyl 6-(1,3-dioxolan-2-yl)hexanoate (prepared according to the procedure described in Syn. Comm. 1991, 1075) for Example 2A in Example 2B. MS (ESI(+)) m/e 189 (M+H)⁺.

Example 220B6-(1,3-dioxolan-2-yl)-N-(4-(4-methoxyphenyl)-1,3-thiazol-2-yl)hexanamide

35 The desired product was prepared by substituting Example 220A and 4-(4'-methoxyphenyl)-2-aminothiazole for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+)) m/e 377 (M+H)⁺.

Example 220CN-(4-(4-methoxyphenyl)-1,3-thiazol-2-yl)-7-oxoheptanamide

A solution of Example 220B (2.06 g, 5.47 mmol) in acetone (40 mL) and water (2 mL) was treated with p-toluenesulfonic acid monohydrate (30 mg), heated to reflux for 48 hours, cooled to room temperature, and diluted with water. The resulting precipitate was collected by filtration and dried to provide 1.3 g of the desired product. MS (ESI(+)) m/e 333 (M+H)⁺.

Example 220Dmethyl 9-((4-(4-methoxyphenyl)-1,3-thiazol-2-yl)amino)-2,9-dioxononanoate

The desired product was prepared by substituting Example 220C for Example 142B in Examples 142C and 142D. MS (ESI(+)) m/e 405 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.15 (s, 1H), 7.82 (d, 2H), 7.42 (s, 1H), 6.98 (d, 2H), 3.79 (s, 3H), 3.76 (s, 3H), 2.81 (t, 2H), 2.43 (t, 2H), 1.62-1.46 (m, 4H), 1.32-1.27 (m, 4H).

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Example 2219-((4-(4-methoxyphenyl)-1,3-thiazol-2-yl)amino)-2,9-dioxononanoic acid

A suspension of Example 220 (96 mg, 0.24 mmol) in acetonitrile (3 mL) and water (1.5 mL) was treated with LiOH (11mg, 0.26 mmol), stirred at room temperature for 30 minutes, diluted with water, and acidified with 1N HCl. The resulting precipitate was collected by filtration to provide 78 mg (85% yield) of the desired product. MS (ESI(+)) m/e 391 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 14.50-13.20 (br s, 1H), 12.15 (s, 1H), 7.81 (d, 2H), 7.42 (s, 1H), 6.98 (d, 2H), 3.79 (s, 3H), 2.75 (t, 2H), 2.43 (t, 2H), 1.64-1.46 (m, 4H), 1.31-1.25 (m, 4H).

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Example 222N-(8-(methylamino)-7,8-dioxooctyl)-3-phenoxybenzamide

The desired product was prepared by substituting 3-phenoxybenzoic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 383 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.53-8.44 (m, 2H), 7.63-7.59 (m, 1H), 7.49-7.39 (m, 4H), 7.19-7.13 (m, 2H), 7.05-7.02 (m, 2H), 3.24-3.18 (m, 2H), 2.79 (t, 2H), 2.64 (d, 3H), 1.54-1.44 (m, 4H), 1.31-1.25 (m, 4H).

Example 223N-(8-(methylamino)-7,8-dioxooctyl)-4-phenoxybenzamide

The desired product was prepared by substituting 4-phenoxybenzoic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS

(ESI(+)) m/e 383 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.53-8.48 (br m, 1H), 8.39-8.35 (m, 1H), 7.85 (d, 2H), 7.46-7.41 (m, 2H), 7.23-7.18 (m, 1H), 7.09-7.06 (m, 2H), 7.02 (d, 2H), 3.23 (q, 2H), 2.79 (t, 2H), 2.64 (d, 3H), 1.55-1.44 (m, 4H), 1.32-1.26 (m, 4H).

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Example 2244-bromo-N-(8-(methylamino)-7,8-dioxooctyl)benzamide

The desired product was prepared by substituting 4-bromobenzoic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 369, 371 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.53-8.47 (m, 2H), 7.77 (d, 2H), 7.66 (d, 2H), 3.26-3.19 (m, 2H), 2.79 (t, 2H), 2.64 (d, 3H), 1.55-1.44 (m, 4H), 1.32-1.26 (m, 4H).

Example 2253-bromo-N-(8-(methylamino)-7,8-dioxooctyl)benzamide

The desired product was prepared by substituting 3-bromobenzoic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 369, 371 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.57-8.47 (m, 2H), 8.01-7.99 (m, 1H), 7.85-7.81 (m, 1H), 7.73-7.70 (m, 1H), 7.43 (t, 1H), 3.27-3.20 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.54-1.45 (m, 4H), 1.33-1.26 (m, 4H).

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Example 226N-(8-(methylamino)-7,8-dioxooctyl)-3-(methylsulfonyl)benzamide

The desired product was prepared by substituting 3-bromobenzoic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. ¹H NMR (DMSO-d₆) δ 8.75-8.71 (br m, 1H), 8.53-8.47 (br m, 1H), 8.36-8.35 (m, 1H), 8.18-8.15 (m, 1H), 8.08-8.05 (m, 1H), 7.75 (t, 1H), 3.31-3.24 (m, 2H), 3.26 (s, 3H), 2.80 (t, 2H), 2.64 (d, 3H), 1.57-1.46 (m, 4H), 1.33-1.28 (m, 4H).

Example 227N-(8-(methylamino)-7,8-dioxooctyl)-4-(1H-pyrrol-1-yl)benzamide

The desired product was prepared by substituting 4-(1H-pyrrol-1-yl)benzoic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. ¹H NMR (DMSO-d₆) δ 8.53-8.42 (m, 2H), 7.92 (d, 2H), 7.68 (d, 2H), 7.48-7.46 (m, 2H), 6.31-6.29 (m, 2H), 3.32-3.24 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.55-1.46 (m, 4H), 1.33-1.29 (m, 4H); Anal. Cald for C₁₇H₂₄N₂O₅S·0.2CF₃COOH: C, 53.42; H, 6.23; N, 7.16. Found: C, 53.05; H, 6.15; N, 6.74.

Example 228

1-methyl-N-(8-(methylamino)-7,8-dioxooctyl)-1H-indole-2-carboxamide

The desired product was prepared by substituting 1-methyl-1H-indole-2-carboxylic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 344 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.53-8.43 (m, 2H), 7.62 (d, 1H), 7.51 (d, 1H), 7.29-7.23 (m, 1H), 7.11-7.06 (m, 1H), 7.04 (s, 1H), 3.97 (s, 3H), 3.27-3.20 (m, 2H), 2.81 (t, 2H), 2.64 (d, 3H), 1.58-1.48 (m, 4H), 1.35-1.29 (m, 4H).

Example 229N-(8-(methylamino)-7,8-dioxooctyl)-2-naphthamide

10 The desired product was prepared by substituting 2-naphthoic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 341 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.61-8.57 (m, 1H), 8.53-8.47 (br m, 1H), 8.43 (s, 1H), 8.03-7.90 (m, 4H), 7.63-7.56 (m, 2H), 3.33-3.27 (m, 2H), 2.81 (t, 2H), 2.64 (d, 3H), 1.60-1.48 (m, 4H), 1.36-1.30 (m, 4H).

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Example 230N-(8-(methylamino)-7,8-dioxooctyl)-1,3-benzodioxole-5-carboxamide

The desired product was prepared by substituting 1,3-benzodioxole-5-carboxylic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 335 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.52-8.47 (br m, 1H), 8.25 (t, 1H), 7.42 (dd, 1H), 7.37 (d, 1H), 6.96 (d, 1H), 6.08 (s, 2H), 3.23-3.17 (m, 2H), 2.79 (t, 2H), 2.64 (d, 3H), 1.54-1.42 (m, 4H), 1.31-1.25 (m, 4H).

Example 231N-(8-(methylamino)-7,8-dioxooctyl)-1-benzofuran-2-carboxamide

25 The desired product was prepared by substituting 1-benzofuran-2-carboxylic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 331 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.68 (t, 1H), 8.52-8.47 (br m, 1H), 7.76 (d, 1H), 7.66-7.63 (m, 1H), 7.51-7.50 (m, 1H), 7.48-7.43 (m, 1H), 7.35-7.30 (m, 1H), 3.29-3.22 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.58-1.46 (m, 4H), 1.34-1.28 (m, 4H).

Example 232N-(8-(methylamino)-7,8-dioxooctyl)-1H-benzimidazole-6-carboxamide

30 The desired product was prepared by substituting 1H-benzimidazole-6-carboxylic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 331 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 9.02 (br s, 1H), 8.59-8.47 (m, 2H),

8.21-8.20 (m, 1H), 7.90 (dd, 1H), 7.77 (d, 1H), 3.31-3.24 (m, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.58-1.47 (m, 4H), 1.34-1.28 (m, 4H).

Example 233

5 N-(8-(methylamino)-7,8-dioxooctyl)-1H-indole-6-carboxamide

The desired product was prepared by substituting 1H-benzimidazole-6-carboxylic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 330 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 11.34 (s, 1H), 8.53-8.48 (br m, 1H), 8.33-8.29 (br m, 1H), 7.92 (s, 1H), 7.57-7.47 (m, 3H), 6.48-6.46 (m, 1H), 3.28-3.22 (m, 2H), 10 2.80 (t, 2H), 2.64 (d, 3H), 1.56-1.47 (m, 4H), 1.35-1.28 (m, 4H).

Example 234

3-chloro-N-(8-(methylamino)-7,8-dioxooctyl)benzamide

The desired product was prepared by substituting 3-chlorobenzoic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 325/327 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.57-8.48 (m, 2H), 7.87-7.86 (m, 1H), 7.81-7.77 (m, 1H), 7.60-7.57 (m, 1H), 7.49 (t, 1H), 3.24 (q, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.55-1.46 (m, 4H), 1.32-1.26 (m, 4H).

20 Example 235

4-methyl-N-(8-(methylamino)-7,8-dioxooctyl)-2-(4-(trifluoromethyl)phenyl)-1,3-thiazole-5-carboxamide

The desired product was prepared by substituting 4-methyl-2-(4-(trifluoromethyl)phenyl)-1,3-thiazole-5-carboxylic acid and Example 142F for Example 1B and 4-aminopyridine, respectively, in Example 6. MS (ESI(+)) m/e 456 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.53-8.47 (br m, 1H), 8.35-8.32 (m, 1H), 8.15 (d, 2H), 7.88 (d, 2H), 3.26-3.19 (m, 2H), 2.81 (t, 2H), 2.64 (d, 3H), 2.62 (s, 3H), 1.56-1.47 (m, 4H), 1.34-1.28 (m, 4H).

Example 236

30 N¹-methyl-2-oxo-N⁸-(4-phenyl-1,3-thiazol-2-yl)octanediamide

Example 236A

8-(methylamino)-7,8-dioxooctanoic acid

The desired product was prepared by substituting methyl 5-hydroxypentanoate for 35 methyl 6-hydroxyhexanoate in Examples 124A through 124E.

Example 236B

N¹-methyl-2-oxo-N⁸-(4-phenyl-1,3-thiazol-2-yl)octanediamide

The desired product was prepared by substituting Example 236A and 4-phenyl-2-aminothiazole for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+)) m/e 369 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.20 (s, 1H), 8.53-8.48 (br m, 1H), 7.89 (d, 2H), 7.59 (s, 1H), 7.45-7.40 (m, 2H), 7.34-7.29 (m, 1H), 2.81 (t, 2H), 2.64 (d, 3H), 2.44 (t, 2H), 1.66-1.47 (m, 4H), 1.35-1.27 (m, 2H).

Example 237N⁹-(4-(2,5-dimethylthien-3-yl)-1,3-thiazol-2-yl)-N¹-methyl-2-oxanonanediamide

10 The desired product was prepared by substituting Example 124E and 4-(2,5-dimethylthien-3-yl)-1,3-thiazol-2-amine for Example 1B and aniline, respectively, in Example 1C. MS (ESI(+)) m/e 408 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 12.06 (s, 1H), 8.52-8.47 (br m, 1H), 7.16 (s, 1H), 7.01 (s, 1H), 2.79 (t, 2H), 2.64 (d, 3H), 2.56 (s, 3H), 2.43 (t, 2H), 2.38 (s, 3H), 1.63-1.45 (m, 4H), 1.32-1.26 (m, 4H).

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Example 238N-methyl-2-oxo-8-(5-thien-2-yl-1,3,4-oxadiazol-2-yl)octanamideExample 238AN-methyl-2,9-dioxo-9-(2-(thien-2-ylcarbonyl)hydrazino)nonanamide

20 A solution of Example 124E (0.1 g, 0.46 mmol) in N,N-dimethylformamide (15 mL) at room temperature was treated with polystyrene supported dicyclohexylcarbodiimide (0.48 g, 0.9 mmol), hydroxybenzotriazole (0.06 g, 0.46 mmol) and thiophene-2-carbohydrazide (0.06 g, 0.46 mmol), stirred for 8 hours, and filtered. The resin was washed with DMF (5 mL) and dichloromethane (5 mL) and the combined washes and filtrate were evaporated to provide the desired product.

Example 238BN-methyl-2-oxo-8-(5-thien-2-yl-1,3,4-oxadiazol-2-yl)octanamide

30 A solution of Example 238A in THF (5 mL) was treated with (methoxycarbonylsulfamoyl)-triethyl ammonium hydroxide (0.22 g, 0.96 mmol), irradiated in a Smith microwave synthesizer at 300 W for 15 minutes, and concentrated. The concentrate was dissolved in 1:1 DMSO:CH₃OH (1.5 mL) and purified by reverse phase HPLC with a gradient of 0 to 95% acetonitrile/water containing 0.1% TFA over 10 minutes to provide the desired product (0.045 g, 31%). MS (ESI(+)) m/e 322 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.5 (s, 1H), 7.90 (d, 1H), 7.78 (d, 1H), 7.30 (t, 1H), 2.90 (t, 2H), 2.80 (t, 2H), 2.60 (d, 3H), 1.78 (m, 2H), 1.50 (m, 2H), 1.2-1.4 (m, 4H).

Example 239N-methyl-2-oxo-8-(5-(3,4,5-trimethoxyphenyl)-1,3,4-oxadiazol-2-yl)octanamide

5 The desired product was prepared by substituting 3,4,5-trimethoxybenzohydrazide for thiophene-2-carbohydrazide in Example 238. MS (ESI(+)) m/e 406 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.5 (s, 1H), 7.20 (s, 2H), 3.90 (s, 6H), 3.70 (s, 3H), 2.90 (t, 2H), 2.80 (t, 2H), 2.60 (d, 3H), 1.78 (m, 2H), 1.50 (m, 2H), 1.2-1.4 (m, 4H).

Example 2409-(2-benzoylhydrazino)-N-methyl-2,9-dioxononanamide

10 The desired product was prepared by substituting benzohydrazide for thiophene-2-carbohydrazide in Example 238A. MS (ESI(+)) m/e 334 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 10.3 (s, 1H), 9.8 (s, 1H), 8.5 (s, 1H), 7.90 (d, 2H), 7.50 (m, 1H), 7.40 (d, 2H), 2.80 (t, 2H), 2.60 (d, 3H), 2.20 (t, 2H), 1.4-1.6 (m, 4H), 1.2-1.4 (m, 4H).

15

Example 2419-(2-(1,1'-biphenyl-4-ylcarbonyl)hydrazino)-N-methyl-2,9-dioxononanamide

20 The desired product was prepared by substituting 1,1'-biphenyl-4-carbohydrazide for thiophene-2-carbohydrazide in Example 238A. MS (ESI(+)) m/e 410 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 10.3 (s, 1H), 9.8 (s, 1H), 8.5 (s, 1H), 8.00 (d, 2H), 7.80 (d, 2H), 7.78 (d, 2H), 7.50 (m, 2H), 7.40 (m, 1H), 2.80 (t, 2H), 2.60 (d, 3H), 2.20 (t, 2H), 1.4-1.6 (m, 4H), 1.2-1.4 (m, 4H).

25

Example 2428-(5-(1,1'-biphenyl-4-yl)-1,3,4-oxadiazol-2-yl)-N-methyl-2-oxooctanamide

The desired product was prepared by substituting Example 241 for Example 238A in Example 238B. MS (ESI(+)) m/e 392 (M+H)⁺; ¹H NMR (DMSO-d₆) δ 8.5 (s, 1H), 8.02 (m, 2H), 7.90 (m, 2H), 7.78 (m, 2H), 7.50 (m, 2H), 7.40 (m, 1H), 2.90 (t, 2H), 2.80 (t, 2H), 2.60 (d, 3H), 1.78 (m, 2H), 1.50 (m, 2H), 1.2-1.4 (m, 4H).

30

Example 2438-(((4-(benzyloxy)phenyl)amino)carbonyl)amino)-N-methyl-2-oxooctanamide

35 The desired product was prepared by substituting 1-isocyanato-4-benzyloxybenzene for isocyanatobenzene in Example 200. MS (ESI(+)) m/e 412 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 8.50 (m, 1H), 8.15 (s, 1H), 7.22-7.45 (m, 7H), 6.84-6.91 (m, 2H), 5.98 (t, 1H), 5.02 (s, 2H), 3.04 (dt, 2H), 2.80 (t, 2H), 2.64 (d, 3H), 1.22-1.55 (m, 8H); Anal. Calcd for C₂₃H₂₉N₃O₄: C, 67.13; H, 7.10; N, 10.21. Found: C, 67.26; H, 7.10; N, 10.11.

Example 2447-(1,1'-biphenyl-4-yloxy)-1-(2H-tetraazol-5-yl)heptan-1-one

5

Example 244A8-(1,1'-biphenyl-4-yloxy)-2-(tetrahydro-2H-pyran-2-yloxy)octanenitrile

A solution of Example 112C (0.92 g, 2.94 mmol) and dihydropyran (0.247 g, 2.94 mmol) in dichloromethane (100 mL) at room temperature was treated with p-toluenesulfonic acid monohydrate (56 mg, 0.29 mmol), stirred for 5 hours, and partitioned between water and dichloromethane. The organic phase was washed with brine, dried (Na_2SO_4), filtered, and concentrated. The concentrate was purified by flash column chromatography on silica gel to provide 1.04 g (90% yield) of the desired product. MS (DCI/NH₃) m/e 393 ($\text{M}+\text{H}$)⁺.

Example 244B

15 5-(7-(1,1'-biphenyl-4-yloxy)-1-(tetrahydro-2H-pyran-2-yloxy)heptyl)-2H-tetraazole
A mixture of example 244B (150 mg, 0.38 mmol), sodium azide (49 mg, 0.76 mmol), and NH₄Cl (41 mg, 0.76 mmol) in DMF (2 mL) was heated to reflux for 5 hours, cooled to room temperature, and partitioned between water and ethyl acetate. The organic phase was washed with water and brine, dried (Na_2SO_4), filtered, and concentrated. The concentrate
20 was purified by flash column chromatography on silica gel with 3% methanol/dichloromethane to provide 80 mg (48% yield) of the desired product. MS (DCI/NH₃) m/e 437 ($\text{M}+\text{H}$)⁺.

Example 244C

25 7-(1,1'-biphenyl-4-yloxy)-1-(2H-tetraazol-5-yl)heptan-1-ol
A solution of Example 244B (60 mg) in methanol (5 mL) at room temperature was treated with p-toluenesulfonic acid monohydrate (10mg), stirred for 3 hours, poured into ice water, and filtered. The filter cake was dried to provide 40 mg (83%) of the desired product. MS (ESI(+)) m/e 353 ($\text{M}+\text{H}$)⁺.

30

Example 244D7-(1,1'-biphenyl-4-yloxy)-1-(2H-tetraazol-5-yl)heptan-1-one

A solution of Example 244C (30 mg) in acetone (2 mL) at room temperature was treated dropwise with Jones reagent until a red-brown color persisted. The reaction mixture
35 was filtered and the filtrate was partitioned between water and ethyl acetate. The organic extract was washed with water, brine, dried (Na_2SO_4), filtered, and concentrated to provide 20 mg (60% yield) of the desired product. MS (DCI/NH₃(+)) m/e 368 ($\text{M}+\text{NH}_4$)⁺; ¹H NMR

(300 MHz, DMSO-d₆) δ 7.70-7.50 (m, 4H), 7.50-7.35 (m, 2H), 7.30-7.20 (m, 1H), 7.05-6.90 (m, 2H), 3.98 (t, 2H), 3.00 (t, 2H), 1.80-1.20 (m, 8H).

Example 245

5 7-(1,1'-biphenyl-3-yloxy)-1-(2H-tetraazol-5-yl)heptan-1-one

Example 245A

8-(1,1'-biphenyl-3-yloxy)-2-hydroxyoctanenitrile

The desired product was prepared by substituting 7-((1,1'-biphenyl)-3-yloxy)heptanal
10 for Example 112B in Example 112C.

Example 245B

7-(1,1'-biphenyl-3-yloxy)-1-(2H-tetraazol-5-yl)heptan-1-one

The desired product was prepared by substituting Example 245A for Example 112C
15 in Example 244. MS (DCI/NH₃(+)) m/e 368 (M+NH₄)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 7.70-7.10 (m, 8H), 7.00-6.90 (m, 1H), 4.02 (t, 2H), 3.08 (t, 2H), 1.80-1.30 (m, 8H).

Example 246

N-1,1'-biphenyl-3-yl-7-oxo-7-(2H-tetraazol-5-yl)heptanamide

20

Example 246A

N-1,1'-biphenyl-3-yl-6-(1,3-dioxolan-2-yl)hexanamide

The desired product was prepared by substituting 1,1'-biphenyl-3-amine and 6-(1,3-dioxolan-2-yl)hexanoic acid (prepared according to the procedure described in Syn. Comm. 1991, 21, 1075) for aniline and Example 1B, respectively, in Example 1C. MS (ESI(+)) m/e 340 (M+H)⁺.
25

Example 246B

N-1,1'-biphenyl-3-yl-7-oxoheptanamide

30 A solution of Example 246A (2.5 g, 7.4 mmol) in acetone (20 mL) and water (10 mL) was treated with p-toluenesulfonic acid monohydrate (70 mg), heated to reflux for 2 days, cooled to room temperature, treated with 2N HCl (2 mL), stirred for 1 hour, and partitioned between water and ethyl acetate. The organic phase was washed with brine, dried (Na₂SO₄), filtered, and concentrated. The concentrate was purified by flash column chromatography on 35 silica gel with 20% ethyl acetate/hexanes to provide 1.5 g of the desired product. ¹H NMR (300 MHz, DMSO-d₆) δ 9.98 (s, 1H), 9.68 (t, 1H), 7.95 (s, 1H), 7.70-7.2 (m, 8H), 2.45 (dt, 2H), 2.34 (t, 2H), 1.7-1.2 (m, 6H).

Example 246CN-1,1'-biphenyl-3-yl-7-cyano-7-hydroxyheptanamide

The desired product was prepared by substituting Example 246B for Example 112B in
5 Example 112C.

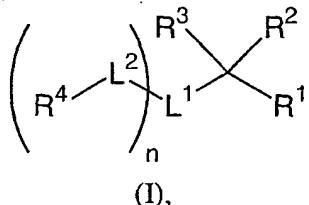
Example 246DN-1,1'-biphenyl-3-yl-7-oxo-7-(2H-tetraazol-5-yl)heptanamide

The desired product was prepared by substituting Example 246C for Example 112C in
10 Example 244. MS (ESI (+)) m/e 364 (M+H)⁺; ¹H NMR (300 MHz, DMSO-d₆) δ 9.98 (s, 1H), 7.96 (t, 1H), 7.70-7.2 (m, 8H), 2.38 (dt, 2H), 2.34 (t, 2H), 1.8-1.3 (m, 6H).

It will be evident to one skilled in the art that the present invention is not limited to
the forgoing illustrative examples, and that it can be embodied in other specific forms without
15 departing from the essential attributes thereof. It is therefore desired that the examples be
considered in all respects as illustrative and not restrictive, reference being made to the
appended claims, rather than to the foregoing examples, and all changes which come within
the meaning and range of equivalency of the claims and therefore intended to be embraced
therein.

WHAT IS CLAIMED IS

1. A compound of formula (I)



5

(I),

or a therapeutically acceptable salt thereof, wherein

n is 1 or 2;

10 L^1 is selected from the group consisting of alkenylene, alkylene, alkynylene, cycloalkylene, heteroalkylene, -(alkylene)-C(O)N(R⁵)-(alkylene)-, -(alkylene)-O-(alkylene)-; wherein each group is drawn with its left-hand end being the end which attaches to L^2 , and its right-hand end being the end which attaches to the carbon substituted with R¹, R², and R³;

15 L^2 is selected from the group consisting of a bond, C₂ alkenylene, -O-, -S-, -SO₂-, -OC(O)NR⁵-, -N(R⁶)C(O)-, -C(O)N(R⁶)-, -SO₂N(R⁶)-, -N(R⁶)SO₂-, -C=N-O-, -N(R⁶)C(O)N(R⁶)-, and -C(O)N(R⁶)N(R⁶)C(O)-; wherein each group is drawn with its left-hand end being the end which attaches to R⁴, and its right-hand end being the end which attaches to L^1 ;

20 R¹ is selected from the group consisting of alkanoyl, alkoxycarbonyl, aminocarbonyl, carboxy, haloalkyl, and heterocycle, wherein the heterocycle is selected from the group consisting of oxazolyl, dihydrooxazolyl, oxadiazolyl, and tetraazolyl;

R² and R³ are hydroxy; or

R² and R³ together are oxo;

25 R⁴ is selected from the group consisting of alkoxyalkyl, alkyl, aryl, arylalkyl, cycloalkyl, (cycloalkyl)alkyl, heterocycle, and (heterocycle)alkyl; and

R⁵ and R⁶ are independently selected from the group consisting of hydrogen, alkyl, aryl, and arylalkyl; or

30 R⁴ and R⁶, together with the nitrogen atom to which they are attached, form a heterocycle selected from the group consisting of morpholinyl, piperazinyl, piperidinyl, and thiomorpholinyl; wherein the morpholinyl, the piperazinyl, the piperidinyl, and the thiomorpholinyl can be optionally substituted with one, two, or three substituents independently selected from the group consisting of alkyl and spiroheterocycle.

2. A compound according to Claim 1 wherein n is 2.

3. A compound according to Claim 1 wherein n is 1.

4. A compound according to Claim 3 wherein R^1 is heterocycle, wherein the heterocycle is selected from the group consisting of oxazolyl, dihydrooxazolyl, oxadiazolyl, and tetraazolyl.
5. A compound according to Claim 4 wherein L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene.
6. A compound according to Claim 3 wherein R^1 is selected from the group consisting of alkoxy carbonyl and carboxy.
7. A compound according to Claim 6 wherein L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene.
8. A compound according to Claim 3 wherein R^1 is alkanoyl.
9. A compound according to Claim 8 wherein L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene.
10. A compound according to Claim 3 wherein R^1 is aminocarbonyl.
11. A compound according to Claim 10 wherein L^1 is -(alkylene)-O-(alkylene)-.
12. A compound according to Claim 10 wherein L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene.
13. A compound according to Claim 12 wherein L^2 is selected from the group consisting of -O-, -S-, -SO₂-, and -SO₂N(R₆)-.
14. A compound according to Claim 12 wherein L^2 is selected from the group consisting of -N(R⁶)C(O)N(R⁶)- and -C(O)N(R⁶)-.
15. A compound according to Claim 12 wherein L^2 is selected from the group consisting of a bond, -C=N-O-, and -N(R⁶)C(O)CHC(O)N(R⁵)(R⁶)-.
16. A compound according to Claim 12 wherein L^2 is -N(R⁶)C(O)-.

17. A compound according to Claim 12 wherein L^2 is selected from the group consisting of $-N(R^6)C(O)N(R^6)-$ and $-C(O)N(R^6)N(R^6)C(O)-$.
18. A compound according to Claim 3 wherein R^1 is haloalkyl.
19. A compound according to Claim 18 wherein L^1 is selected from the group consisting of alkenylene, wherein the alkenylene is C_6 alkenylene; alkynylene, wherein the alkynylene is C_6 alkynylene; cycloalkylene; and $-(alkylene)C(O)N(R^5)(alkylene)-$.
20. A compound according to Claim 18 wherein L^1 is alkylene, wherein the alkylene is C_5 - C_7 alkylene.
21. A compound according to Claim 20 wherein L^2 is C_2 alkenylene.
22. A compound according to Claim 20 wherein L^2 is $-OC(O)N(R^5)-$.
23. A compound according to Claim 20 wherein L^2 is $-O-$.
24. A compound according to Claim 20 wherein L^2 is $-N(R^6)C(O)-$.
25. A compound according to Claim 24 wherein R^4 is selected from the group consisting of alkoxyalkyl and alkyl.
26. A compound according to Claim 24 wherein R^4 is aryl.
27. A compound according to Claim 24 wherein R^4 is arylalkyl.
28. A compound according to Claim 24 wherein R^4 is selected from the group consisting of cycloalkyl, heterocycle, and (heterocycle)alkyl.
29. A compound according to Claim 24 wherein R^4 and R^6 , together with the nitrogen atom to which they are attached, form a ring selected from the group consisting of morpholinyl, piperazinyl, piperidinyl, and thiomorpholinyl.
30. A pharmaceutical composition comprising a compound of Claim 1, or a therapeutically acceptable salt thereof, in combination with a pharmaceutically acceptable carrier.

31. A method of inhibiting histone deacetylase in a patient in recognized need of such treatment comprising administering to the patient a therapeutically acceptable amount of a compound of Claim 1, or a therapeutically acceptable salt thereof.

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(54) Title: INHIBITORS OF HISTONE DEACETYLASE

(57) Abstract: Compounds having the formula or therapeutically acceptable salts thereof, are histone deacetylase (HDAC) inhibitors. Preparation of the compounds, compositions containing the compounds, and treatment of diseases using the compounds are disclosed.

INTERNATIONAL SEARCH REPORT

Int'l Application No

PCT/US 01/50931

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07C235/14 C07C235/16 C07C235/20 C07C235/74 C07C235/80
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Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07C C07D A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, CHEM ABS Data, BEILSTEIN Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>MANFRED JUNG ET AL: "Amide Analogues of Trichostatin A as Inhibitors of Histone Deacetylase and Inducers of Terminal Cell Differentiation" JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY, WASHINGTON, US, vol. 42, no. 22, 4 November 1999 (1999-11-04), pages 4669-4679, XP002144226 ISSN: 0022-2623 page 4669, column 1, line 1 -page 4671, column 2, line 30</p> <p>---</p> <p style="text-align: center;">-/-</p>	1,30,31

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
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T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

& document member of the same patent family

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INTERNATIONAL SEARCH REPORT

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	SUZUKI ET AL: "Synthesis and Histone Deacetylase Inhibitory Activity of New Benzamide Derivatives" JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY, WASHINGTON, US, vol. 42, no. 15, 1999, pages 3001-3003, XP002158227 ISSN: 0022-2623 the whole document -----	1,30,31

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 01/50931

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
Although claim 31 is directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. Claims Nos.: 1 – 31 (all in part) because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210
3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.
 No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 1 - 31 (all in part)

Present claims 1 to 31 relate to an extremely large number of possible compounds. Support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT is to be found, however, for only a very small proportion of the compounds/products/apparatus/methods claimed. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Consequently, the search has been carried out for those parts of the claims which appear to be supported and disclosed, namely those parts relating to the compounds prepared in the examples.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

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